



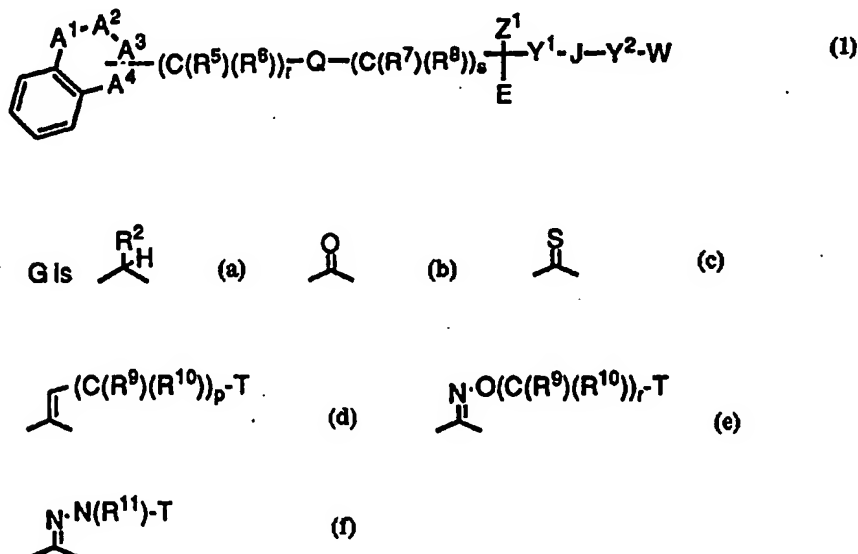
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(21) International Application Number: PCT/US96/07960 (22) International Filing Date: 4 June 1996 (04.06.96) (30) Priority Data: 08/469,315 6 June 1995 (06.06.95) US (71) Applicant: SCHERING CORPORATION [US/US]; 2000 Galloping Hill Road, Kenilworth, NJ 07033 (US). (72) Inventors: McCORMICK, Kevin, D.; 5 Pace Drive, Edison, NJ 08820 (US). LUPO, Andrew, T., Jr.; 8 Powell Road, Emerson, NJ 07630 (US). (74) Agents: MAGATTI, Anita, W. et al.; Schering-Plough Corporation, Patent Dept. K-6-1 1990, 2000 Galloping Hill Road, Kenilworth, NJ 07033-0530 (US).	(81) Designated States: AL, AM, AU, AZ, BB, BG, BR, BY, CA, CN, CZ, EE, GE, HU, IL, IS, JP, KG, KR, KZ, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>	

(54) Title: **SUBSTITUTED BENZENE-FUSED HETERO- AND CARBOCYCLICS AS NEUROKININ ANTAGONISTS**

(57) Abstract

Compounds represented by structural formula (1), or a pharmaceutically acceptable salt thereof, wherein: A¹, A², A³ and A⁴ are methylene, -CH= or -NH-, all optionally substituted, or -N=, -O-, -S(O)_e, -C(O)- or a bond; E is phenyl or heteroaryl; W is cycloalkyl, aryl, heterocycloalkyl or heteroaryl; R⁵, R⁷, R⁹, R¹¹, R¹² and R¹⁴ are H, alkyl, phenyl, benzyl, -CF₃ or -C₂F₅; R¹⁰ is H or alkyl; R⁶ and R⁸ are R⁵, or optionally substituted alkyl further substituted by -OR¹¹, -NR¹¹R¹², -SH, -S(O)_eR¹³, -CO₂R¹¹, -OC(O)R¹¹, -CONR¹¹R¹², -COR¹¹ or -NR¹¹C(O)R¹²; R¹³ is alkyl, phenyl, benzyl, -CF₃ or -C₂F₅; Q is a bond, -C(O)-, -NR¹⁷, -(C(R⁹)(R¹⁰))_r, -O-, -S(O)_e, -C(X)NR¹¹, -N(R¹¹)C(X)-, -N(R¹¹)SO₂- or -SO₂N(R¹¹)-; R¹⁷ is H, alkyl, -S(O)_eR¹³, -COR¹¹, -CO₂R¹³, -CONR¹¹R¹², allyl, -(C₁-C₆ alkyl)-CH=CH₂, phenyl or benzyl; X is -O-, -S or -N(R¹²); Y¹ is -(C(R⁹)(R¹⁰))_m, -G-(C(R⁹)(R¹⁰))_m- or -(C(R⁹)(R¹⁰))_m-G-; G represents formulas (a); (b); (c); (d); (e) or (f), with certain provisos; T is H, aryl, heterocycloalkyl, heteroaryl, cycloalkyl, -OR¹¹, -N(R¹¹)(R¹²), -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), -S(O)_eR¹³, -NR¹¹CO₂R¹³, -NR¹¹COR¹², -NR¹¹CON(R¹²)(R¹⁴) or -OC(O)N(R¹¹)(R¹²); V is a bond, -S(O)_e, -O- or -N(Z²)-, -N(Z²)C(O)- or -N(Z²)C(S)-; Y² is -(C(R⁹)(R¹⁰))_n; Z¹ is Z², -NR¹¹R¹², -OR¹¹ or SR¹¹; Z² is H, alkyl, phenyl, benzyl, -CF₃ or -C₂F₅; or Z¹ and Z² together are -(C(R⁹)(R¹⁰))_u, wherein u is 1-4; e and n are 0-2; m and p are 0-3; and r and s are 1-4; wherein cycloalkyl, heterocycloalkyl, phenyl, benzyl, aryl and heteroaryl groups are all optionally substituted; methods of treating asthma, cough, bronchospasm, inflammatory diseases, and gastrointestinal disorders with said compounds, and pharmaceutical compositions comprising said compounds are disclosed.



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**SUBSTITUTED BENZENE-FUSED HETERO- AND
CARBOCYCLICS AS NEUROKININ ANTAGONISTS**

BACKGROUND OF THE INVENTION

The present invention relates to a genus of substituted benzene-fused hetero- and carbocyclics useful as antagonists of tachykinin receptors, in particular as antagonists of the neuropeptides neurokinin-1 receptor (NK₁) and/or neurokinin-2 receptor (NK₂) and/or neurokinin-3 receptor (NK₃).

Neurokinin receptors are found in the nervous system and the circulatory system and peripheral tissues of mammals, and therefore are involved in a variety of biological processes. Neurokinin receptor antagonists are consequently expected to be useful in the treatment or prevention of various mammalian disease states, for example asthma, cough, bronchospasm, inflammatory diseases such as arthritis, central nervous system conditions such as migraine and epilepsy, nociception, and various gastrointestinal disorders such as Crohn's disease.

In particular, NK₁ receptors have been reported to be involved in microvascular leakage and mucus secretion, and NK₂ receptors have been associated with smooth muscle contraction, making NK₁ and NK₂ receptor antagonists especially useful in the treatment and prevention of asthma.

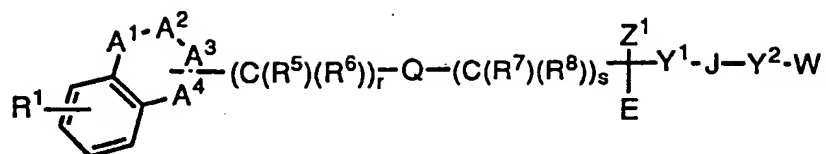
Some NK₁ and NK₂ receptor antagonists have previously been disclosed: arylalkylamines were disclosed in U.S. Patent 5,350,852, issued September 27, 1994, and spiro-substituted azacycles were disclosed in WO 94/29309, published December 22, 1994.

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SUMMARY OF THE INVENTION

Compounds of the present invention are represented by the formula I

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or a pharmaceutically acceptable salt thereof, wherein:

A¹, A², A³ and A⁴ are independently selected from the group consisting of -(C(R²)(R¹⁰))- , -(C(R^{2c})(R¹⁰))- , -(C(R²))= , -(C(R^{2c}))= , -NR¹⁶- ,
 5 -N= , -O- , -S(O)_e- , -C(O)- and a bond, wherein A¹, A², A³ and A⁴, together with the carbon atoms to which they are attached, form a 5- or 6-membered ring, and wherein two adjacent A groups are selected from the group consisting of the following combinations:

- 10 -(C(R²)(R¹⁰))-(C(R²)(R¹⁰))-;
- (C(R²)(R¹⁰))-(C(R^{2c}))=;
- (C(R^{2c})(R¹⁰))-NR¹⁶-;
- (C(R^{2c})(R¹⁰))-N=;
- (C(R^{2c})(R¹⁰))-O-;
- (C(R^{2c})(R¹⁰))-S(O)_e-;
- 15 -(C(R²)(R¹⁰))-C(O)-;
- (C(R²))=(C(R²))-;
- (C(R²))=N-;
- =(C(R²))-(C(R²))=;
- =(C(R²))-NR¹⁶-;
- 20 =(C(R²))-N=;
- =(C(R²))-O-;
- =(C(R²))-S(O)_e-;
- =(C(R²))-C(O)-;
- NR¹⁶-N=;
- 25 -NR¹⁶-S(O)_e-;
- NR¹⁶-C(O)-;
- N=N-;
- =N-N=;
- =N-O-;
- 30 =N-S(O)_e-;
- =N-C(O)-;
- O-S(O)_e-; and
- O-C(O)-;

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provided that three adjacent A groups do not represent -C(O)-O-C(O)-, -S(O)-O-C(O)- or -S(O)-O-S(O)-, and provided that when an aromatic nitrogen is present in the ring formed by A¹, A², A³ and A⁴, the N-oxide can be formed;

5 E is R³-aryl or R³-heteroaryl;

W is R⁴-cycloalkyl, R⁴-aryl, R⁴-heterocycloalkyl or R⁴-heteroaryl;

R¹, R³ and R⁴ are independently 1-3 substituents independently selected from the group consisting of H, halogeno, C₁-C₆ alkyl, -CF₃, -C₂F₅, -OR¹¹, -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²),
10 -N(R¹¹)(R¹²), -N(R¹¹)COR¹², -SH, -S(O)_eR¹³, -OC(O)R¹¹, -OC(O)N(R¹¹)(R¹²), -NR¹¹CO₂R¹³, -NR¹¹C(O)N(R¹²)(R¹⁴), R¹⁵-phenyl, R¹⁵-benzyl, -NO₂, -NR¹¹S(O)₂R¹³ and -S(O)₂NR¹¹R¹²; or adjacent R¹, R³ or R⁴ substituents can form a -O-CH₂-O- group;

15 R² is independently selected from the group consisting of R^{2a}, R^{2b} or R¹⁰; and R^{2c} is independently selected from the group consisting of R^{2a} and R¹⁰; wherein R^{2a} is selected from the group consisting of -CF₃, -C₂F₅, -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), R¹⁵-phenyl and R¹⁵-benzyl; and R^{2b} is selected from the group consisting of
20 halogeno, -OR¹¹, -NO₂, -N(R¹¹)(R¹²), -N(R¹¹)COR¹², -SH, -S(O)_eR¹³, -OC(O)R¹¹, -OC(O)N(R¹¹)(R¹²), -NR¹¹CO₂R¹³, -NR¹¹C(O)N(R¹²)(R¹⁴), -NR¹¹S(O)₂R¹³ and -S(O)₂NR¹¹R¹²; provided that from any combination of ring members A¹, A², A³ and A⁴ comprising R² and R^{2c}, no more than one substituent can be selected from R^{2a} and no more than one
25 substituent can be selected from R^{2b};

R⁵, R⁷, R⁹, R¹¹, R¹² and R¹⁴ are independently selected from the group consisting of H, C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃ and -C₂F₅;

R⁶ and R⁸ are independently selected from the group consisting of R⁵, -(C(R⁹)(R¹⁰))_n-OR¹¹, -(C(R⁹)(R¹⁰))_n-NR¹¹R¹²,
30 -(C(R⁹)(R¹⁰))_n-SH, -(C(R⁹)(R¹⁰))_n-S(O)_eR¹³, -(C(R⁹)(R¹⁰))_n-CO₂R¹¹, -(C(R⁹)(R¹⁰))_n-OC(O)R¹¹, -(C(R⁹)(R¹⁰))_n-CONR¹¹R¹², -(C(R⁹)(R¹⁰))_n-COR¹¹ and -(C(R⁹)(R¹⁰))_n-NR¹¹C(O)R¹², provided that when Q is a heteroatom, R⁶ and R⁸ cannot be -OR¹¹, -OC(O)R¹¹,
35 -N(R¹¹)COR¹², -NR¹¹R¹², -SH or -S(O)_eR¹³ on adjacent carbon atoms;

R¹⁰ is independently selected from the group consisting of hydrogen and C₁-C₆ alkyl;

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R¹³ is independently selected from the group consisting of C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃ and -C₂F₅;

R¹⁵ is hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy or halogeno;

Q is a bond, -C(O)-, -NR¹⁷-, -(C(R⁹)(R¹⁰))-, -O-, -S(O)_e-,
 5 -C(X)NR¹¹-, -N(R¹¹)C(X)-, -N(R¹¹)SO₂-, -SO₂N(R¹¹)- or -N⁺(R¹¹)(R¹⁷)-;

R¹⁶ and R¹⁷ are independently selected from the group consisting of hydrogen, C₁-C₆ alkyl, -S(O)_eR¹³, -COR¹¹, -(CH₂)_m-CO₂R¹³, -CONR¹¹R¹², alkenyl, R¹⁵-phenyl and R¹⁵-benzyl;

X is =O, =S or =N(R¹²);

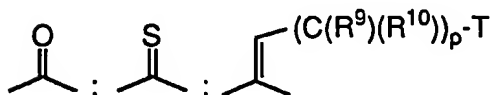
10 Y¹ is -(C(R⁹)(R¹⁰))_m-, -G-(C(R⁹)(R¹⁰))_m- or -(C(R⁹)(R¹⁰))_m-G-;

G is



, provided that when m is 0, R² is H, C₁-C₆ alkyl, -CF₃, -C₂F₅,

-COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), R¹⁵-phenyl or R¹⁵-benzyl;



, provided that when p is 0, T is

not OH or -NR¹¹R¹²; , provided that when r is 1,

T is not OR¹¹ or -NR¹¹R¹²; or , provided T is not

-OR¹¹, -N(R¹¹)(R¹²), -S(O)_eR¹³, -NR¹¹CO₂R¹³, -NR¹¹COR¹²,
 -NR¹¹CON(R¹²)(R¹⁴) or -OC(O)N(R¹¹)(R¹²);

15 T is H, R¹⁵-aryl, R¹⁵-heterocycloalkyl, R¹⁵-heteroaryl, R¹⁵-cycloalkyl, -OR¹¹, -N(R¹¹)(R¹²), -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²),
 -S(O)_eR¹³, -NR¹¹CO₂R¹³, -NR¹¹COR¹², -NR¹¹CON(R¹²)(R¹⁴) or
 -OC(O)N(R¹¹)(R¹²);

20 J is a bond, -S(O)_e-, -O- or -N(Z²)-, -N(Z²)C(O)- or
 -N(Z²)C(S)-; and when G is -C(R²)H-, J can also be -N(Z²)C(O)O- or
 -OC(O)N(Z²)-;

Y² is -(C(R⁹)(R¹⁰))_m-;

25 Z¹ is H, C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃, -C₂F₅,
 -NR¹¹R¹², -OR¹¹ or SR¹¹; Z² is H, C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl,
 -CF₃ or -C₂F₅; provided that when Y¹ is -(C(R⁹)(R¹⁰))_m- and m is 0, Z¹ is
 not -NR¹¹R¹², -OR¹¹ or -SR¹¹; or Z¹ and Z² together are -(C(R⁹)(R¹⁰))_u-;

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wherein u is 1 to 4, and wherein with the atoms to which they are attached, form a 4 to 8 membered ring;

e and n are independently 0, 1 or 2;

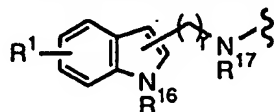
m and p are independently 0, 1, 2 or 3; and

5 r and s are independently 1, 2, 3 or 4.

Preferred are compounds of formula I wherein W is R⁴-cycloalkyl, R⁴-aryl or R⁴-heteroaryl. Preferred R¹, R², R³ and R⁴ substituents are H, halogeno, C₁-C₆ alkyl, -CF₃, -OR¹¹, -COR¹¹, -CO₂R¹¹,
 10 -CON(R¹¹)(R¹²) and -N(R¹¹)(R¹²). R⁵, R⁷, R⁹, R¹¹, R¹² and R¹⁴ are preferably independently H, C₁-C₆ alkyl or -CF₃. R⁶ and R⁸ are preferably independently hydrogen, -(C(R⁹)(R¹⁰))_n-OR¹¹, -(C(R⁹)(R¹⁰))_n-NR¹¹R¹², -(C(R⁹)(R¹⁰))_n-SH-, (C(R⁹)(R¹⁰))_n-S(O)_eR¹³, -(C(R⁹)(R¹⁰))_n-CO₂R¹¹, -(C(R⁹)(R¹⁰))_n-CONR¹¹R¹² or -(C(R⁹)(R¹⁰))_n-COR¹¹; more preferably, R⁶
 15 and R⁸ are each hydrogen or R⁸ is hydrogen and R⁶ is -(C(R⁹)(R¹⁰))_n-CO₂R¹¹. R¹³ is preferably C₁-C₆ alkyl or R¹⁵-phenyl. Q is preferably -NR¹⁷-, -O- or -S(O)_e-, with -NR¹⁷- being more preferred. When Q is -NR¹⁷-, R¹⁷ is preferably hydrogen, C₁-C₆ alkyl or C₂-C₆ alkenyl.

E is preferably R³-phenyl. Z¹ is preferably hydrogen, or Z¹
 20 and Z² together are ethylene or propylene, and with the atoms to which they are attached form a 5- or 6-membered ring. Y¹ is preferably -(C(R⁹)(R¹⁰))_m or -(C(R⁹)(R¹⁰))_m-G, wherein m is preferably 0 or 1. In the definition of Y², m is preferably 0 or 1. J is preferably -O-, -N(Z²)- or -N(Z²)C(O)-.

25 Preferred are compounds wherein A⁴ is a bond. Also preferred are compounds wherein A¹, A², A³ and A⁴ comprise an indolyl ring. Preferred compounds of formula I comprise compounds wherein A¹, A², A³ and A⁴ comprise an indolyl ring, R⁵ and R⁶ are each hydrogen and Q is -NR¹⁷-, that is, compounds having the partial structure



30 Also preferred are compounds of formula I wherein R⁷ and R⁸ are each hydrogen, s is 2, Y¹ is -(C(R⁹)(R¹⁰))_m- or -(C(R⁹)(R¹⁰))_m-G-, and J is -O-, -N(Z²)- or -N(Z²)C(O)-. Another group of preferred compounds is that wherein Y¹ is -(C(R⁹)(R¹⁰))_m- and J is -N(Z²)C(O)-.

35 This invention also relates to the use of a compound of formula I in the treatment of asthma, cough, bronchospasm, inflammatory

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diseases such as arthritis, central nervous system conditions such as migraine and epilepsy, nociception, and various gastrointestinal disorders such as Crohn's disease.

5 In another aspect, the invention relates to a pharmaceutical composition comprising a compound of formula I in a pharmaceutically acceptable carrier. The invention also relates to the use of said pharmaceutical composition in the treatment of asthma, cough, bronchospasm, inflammatory diseases such as arthritis, migraine, nociception, and various gastrointestinal disorders such as Crohn's
10 disease.

DETAILED DESCRIPTION

As used herein, the term "alkyl" means straight or branched alkyl chains. "Lower alkyl" refers to alkyl chains of 1-6 carbon atoms and,
15 similarly, lower alkoxy refers to alkoxy chains of 1-6 carbon atoms.

"Alkenyl" means a straight or branched alkane chain of 2-6 carbon atoms having one double bond.

"Cycloalkyl" refers to cyclic alkyl groups of 3-6 carbon atoms.

"Aryl" means phenyl, naphthyl, indenyl, tetrahydronaphthyl, indanyl, anthracenyl or fluorenyl.
20

"Halogeno" refers to fluoro, chloro, bromo or iodo atoms.

"Heterocycloalkyl" refers to 4- to 6-membered saturated rings comprising 1 to 3 heteroatoms independently selected from the group consisting of -O-, -S- and -N(R¹⁶)-, with the remaining ring members being
25 carbon. Examples of heterocycloalkyl rings are tetrahydrofuranyl, pyrrolidinyl, piperidinyl, morpholinyl, thiomorpholinyl and piperazinyl. R⁴-heterocycloalkyl refers to such groups wherein substitutable ring carbon atoms have an R⁴ substituent.

"Heteroaryl" refers to 5- to 10-membered single or
30 benzofused aromatic rings comprising 1 to 4 heteroatoms independently selected from the group consisting of -O-, -S- and -N=, provided that the rings do not include adjacent oxygen and/or sulfur atoms. Examples of single-ring heteroaryl groups are pyridyl, isoxazolyl, oxadiazolyl, furanyl, pyrrolyl, thienyl, imidazolyl, pyrazolyl, tetrazolyl, thiazolyl, thiadiazolyl,
35 pyrazinyl, pyrimidinyl, pyridazinyl and triazolyl. Examples of benzofused heteroaryl groups are indolyl, quinolinyl, thianaphthenyl and benzofurazanyl. N-oxides of nitrogen-containing heteroaryl groups are also included. All positional isomers are contemplated, e.g., 1-pyridyl, 2-

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pyridyl, 3-pyridyl and 4-pyridyl. R⁴-heteroaryl refers to such groups wherein substitutable ring carbon atoms have an R⁴ substituent.

Those skilled in the art will appreciate that the groups comprising A¹, A², A³ and A⁴ are shown in the above description without a free valence, but that one of A¹, A², A³ or A⁴ must have a free valence to bond to the -(C(R⁵)(R⁶))_r group, that is, one of the R², R^{2c}, R¹⁰ or R¹⁶ substituents on one of A¹, A², A³ or A⁴ is replaced by a bond to the -(C(R⁵)(R⁶))_r group.

The proviso in the definitions of R⁶ and R⁸ relating to compounds wherein Q is a heteroatom is intended to allow for -OR¹¹, -OC(O)R¹¹, -N(R¹¹)COR¹², -NR¹¹R¹², -SH or -S(O)_eR¹³ substitution on compounds wherein r and/or s is 2 or 3, as long as the carbon adjacent to Q is not substituted by those groups. The proviso that three adjacent A groups do not represent -C(O)-O-C(O)-, -S(O)-O-C(O)- or -S(O)-O-S(O)- is intended to eliminate unstable ring systems.

In the definition of Q, the group -N⁺(R¹¹)(R¹⁷)- refers to quaternary amine groups.

In the above definitions, wherein R⁵, R⁷, R⁹, R¹¹, R¹² and R¹⁴ are said to be independently selected from a group of substituents, we mean that R⁵, R⁷, R⁹, R¹¹, R¹² and R¹⁴ are independently selected, but also that where an R⁵, R⁷, R⁹, R¹¹, R¹² or R¹⁴ variable occurs more than once in a molecule, those occurrences are independently selected. Similarly, R¹, R³, R⁴ and the R² variables can be independently selected from a group of substituents, and where more than one of those variables is present, the substituents are independently selected; those skilled in the art will recognize that the size and nature of the substituent(s) will affect the number of substituents which can be present.

Compounds of the invention can have at least one asymmetrical carbon atom and therefore all isomers, including diastereomers, enantiomers and rotational isomers are contemplated as being part of this invention. The invention includes d and l isomers in both pure form and in admixture, including racemic mixtures. Isomers can be prepared using conventional techniques, either by reacting optically pure or optically enriched starting materials or by separating isomers of a compound of formula I.

Those skilled in the art will appreciate that for some compounds of formula I, one isomer will show greater pharmacological activity than other isomers.

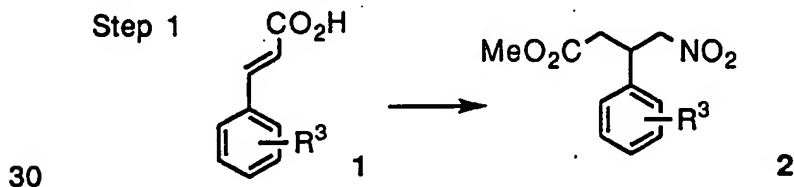
Compounds of the invention have at least one amino group which can form pharmaceutically acceptable salts with organic and inorganic acids. Examples of suitable acids for salt formation are hydrochloric, sulfuric, phosphoric, acetic, citric, oxalic, malonic, salicylic, malic, fumaric, succinic, ascorbic, maleic, methanesulfonic and other mineral and carboxylic acids well known to those in the art. The salt is prepared by contacting the free base form with a sufficient amount of the desired acid to produce a salt. The free base form may be regenerated by treating the salt with a suitable dilute aqueous base solution such as dilute aqueous sodium bicarbonate. The free base form differs from its respective salt form somewhat in certain physical properties, such as solubility in polar solvents, but the salt is otherwise equivalent to its respective free base forms for purposes of the invention.

Certain compounds of the invention are acidic (e.g., those compounds which possess a carboxyl group). These compounds form pharmaceutically acceptable salts with inorganic and organic bases. Examples of such salts are the sodium, potassium, calcium, aluminum, gold and silver salts. Also included are salts formed with pharmaceutically acceptable amines such as ammonia, alkyl amines, hydroxyalkylamines, N-methylglucamine and the like.

Compounds of formula I can be prepared using methods well known to those skilled in the art. Following are typical procedures for preparing various compounds; the skilled artisan will recognize that other procedures may be applicable, and that the procedures may be suitable modified to prepare other compounds within the scope of formula I.

Procedure A:

Compounds of formula I wherein $-Q-(C(R^7)(R^8))_5-$ is $-N(R^{11})CH_2CH_2-$, Z^1 is a hydrogen, Y^1 is $-CH_2-$, and J is $-N(CH_3)C(O)-$ can be prepared as shown in the following reaction scheme:

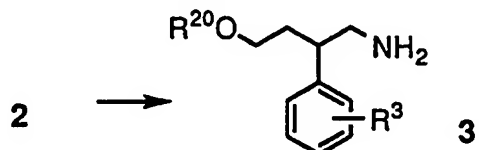


In step 1, a 3-(R^3 -phenyl)-2-propenoic acid (1) is treated with a lower alkyl alcohol such as CH_3OH in the presence of a suitable acid catalyst such as HCl in the range of 0 to $100^\circ C$ to give the corresponding ester. This ester is reacted with a nitroalkane such as CH_3NO_2 in the

- 9 -

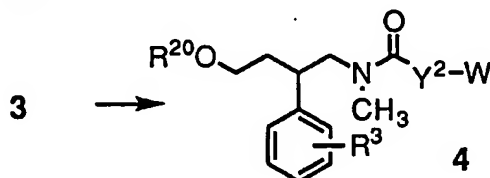
presence of a suitable base such as benzyltrimethylammonium hydroxide at a temperature range of 0 to 100°C to give the desired compound 2.

Step 2



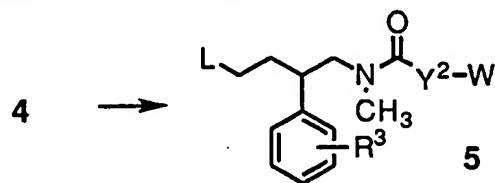
In step 2, compound 2 is reacted with a strong reducing agent such as $LiAlH_4$ or $BH_3 \cdot DMS$ in an inert organic solvent such as THF, ether or benzene, preferably THF, at a temperature range from 0 to 80°C. The resulting amino alcohol is reacted with an electrophile such as a compound of formula $R^{20}L^1$, wherein R^{20} is a suitable hydroxyl-protecting group such as $(R^{21})_3Si-$, wherein R^{21} are independently selected lower alkyl groups (e.g., methyl, ethyl, isopropyl or *t*-butyl), and L^1 is a leaving group such as Cl or Br.

Step 3



In step 3, amine 3 is acylated by standard procedures, for example by treatment with an acid chloride, WY^2COCl , in the presence of an amine base in an inert organic solvent such as CH_2Cl_2 or toluene, preferably CH_2Cl_2 , at a temperature of from -10 to 50°C. Suitable bases include $(CH_3)_3N$, Et_3N and pyridine, preferably Et_3N . Other coupling methods known to those skilled in the art, such as EDC coupling, may also be employed. The resulting amide is treated with a base such as NaH or LDA, in an inert organic solvent such as THF, ether, DMSO or DMF, preferably THF. The resulting anion is treated with an alkylating agent $R^{11}L$, wherein R^{11} is as defined above and L is a suitable leaving group such as Cl, Br, I, triflate or mesylate; preferably $R^{11}L$ is CH_3I . The reactions are typically run at 0 to 50°C.

Step 4

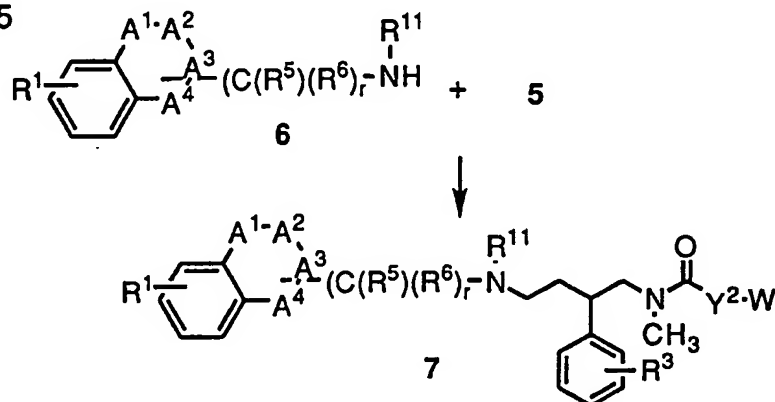


In step 4, the free hydroxyl group is regenerated by treatment of compound 4 with a fluoride reagent such as tetrabutylammonium

- 10 -

fluoride. The hydroxyl group is converted to a suitable leaving group L, as defined above, preferably mesylate. The corresponding mesylate can be obtained by treatment with $\text{CH}_3\text{SO}_2\text{Cl}$ in a suitable solvent such as CH_2Cl_2 in the presence of a base such as Et_3N .

Step 5

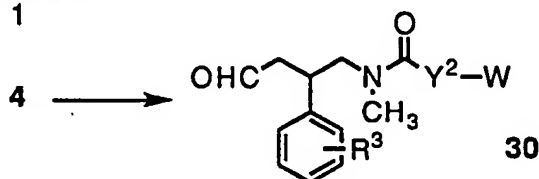


5

In step 5, compound 5 is treated with compound 6 in an inert solvent such as CH_2Cl_2 , THF or DMF, preferably DMF, preferably with a catalytic amount of NaI , and preferably at temperatures from 20 to 80°C .

Procedure A1:

Step 1



10

Alternatively to Procedure A, compound 4 is treated with a fluoride reagent such as tetrabutylammonium fluoride to regenerate the free hydroxyl group. The resulting compound is converted to aldehyde 30 by a suitable oxidation procedure, for example by the Swern procedure as described in *Tetrahedron*, 1978, 34, 1651.

15

Step 2



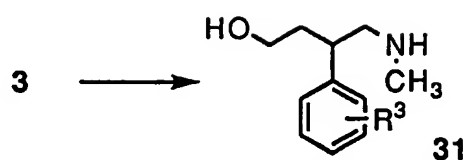
In step 2, compound 30 is reacted with an amine of formula 6 in an alcohol such as CH_3 , $\text{CH}_3\text{CH}_2\text{OH}$ or more preferably $\text{CF}_3\text{CH}_2\text{OH}$, in the presence of a dehydrating agent such as molecular sieves and a reducing agent such as NaBH_3CN or under hydrogenation conditions ($\text{H}_2/\text{Pd/C}$). A suitable temperature range is 0 to 60°C .

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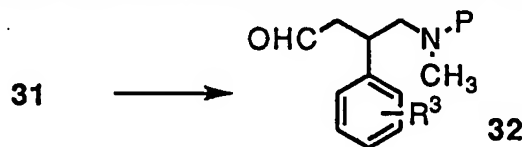
Procedure A2:

Step 1



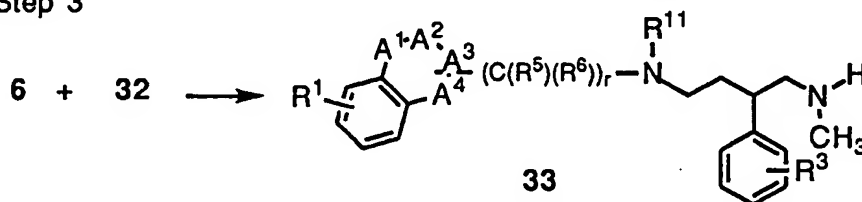
As another alternative to Procedure A, compound 3 is converted to the formamide by standard procedures, for example by treatment with ethylformate, preferably at a temperature from 30 to 60°C. The resulting formamide is treated with a suitable reducing reagent such as BH₃·DMS or AlH₃, preferably BH₃·DMS. The resulting amine-borane complex and the silyl protecting group are hydrolyzed by treatment with aqueous acid such as HCl, preferably at a temperature from 50 to 100°C.

Step 2



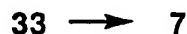
In step 2, the amino group in compound 31 is suitably protected such as by treatment with di-*t*-butyl dicarbonate to obtain the *t*-butyl carbamate. The hydroxyl group is converted to the aldehyde by a suitable oxidation procedure, for example by the Swern procedure.

Step 3



In step 3, compound 32 is reacted with an amine of formula 6 in manner such as that described in Procedure A1, Step 2. The free methylamino group is regenerated by treatment with acid such as trifluoroacetic acid or HCl in the presence of a suitable solvent such as CH₂Cl₂ or CH₃OH.

Step 4



In step 4, compound 33 is acylated by standard procedures, for example by treatment with an acid chloride, WY₂COCl, in the presence of an amine base in an inert organic solvent such as CH₂Cl₂ or toluene, preferably CH₂Cl₂, at a temperature of from -10 to 50°C. Suitable bases include (CH₃)₃N, Et₃N and pyridine, preferably Et₃N. Other coupling

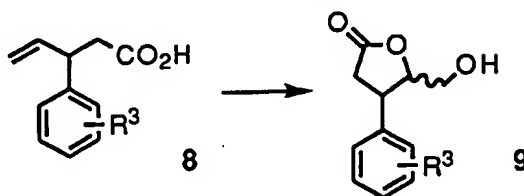
- 12 -

methods known to those skilled in the art, such as EDC coupling, may also be employed.

Procedure B:

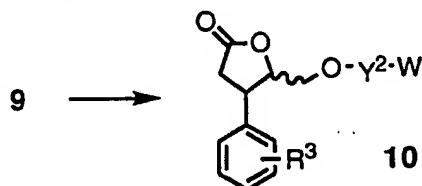
- Compounds of formula I wherein $-Q-(C(R^7)(R^8))_5-$ is $-N(R^{11})CH_2CH_2-$, Z^1 is hydrogen, Y^1 is $-C(OH)H-$, and J is $-O-$ can be prepared as shown in the following reaction scheme:

Step 1



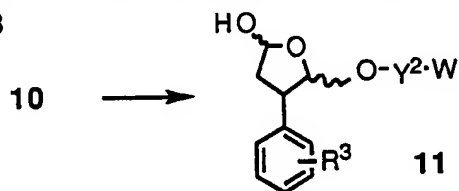
- In step 1, the 3-(R^3 -phenyl)-4-pentenoic acid **8**, is reacted with an oxidizing agent such as dimethyl dioxirane or *m*-CPBA in an inert organic solvent such as CH_2Cl_2 or toluene, preferably at reaction temperatures from 0 to 60°C. An acidic catalyst such as Amberlyst-15 or formic acid is added to give the desired lactone **9**.

Step 2



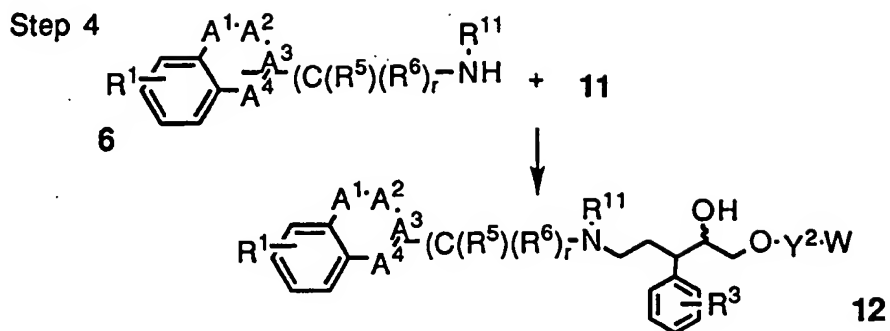
- In step 2, lactone **10** is reacted with an electrophile L^1-Y^2-W wherein L^1 is a leaving group such as Cl or Br. The reaction is carried out in the presence of a silver salt such as Ag_2O in an organic solvent such as DMF or THF, most preferably DMF, at a temperature of 0 to about 50°C.

Step 3



- In step 3, compound **10** is dissolved in an inert organic solvent such as CH_2Cl_2 , THF or toluene, preferably CH_2Cl_2 , and treated with a reducing agent such as DIBAL-H at temperatures from about -78°C to room temperature.

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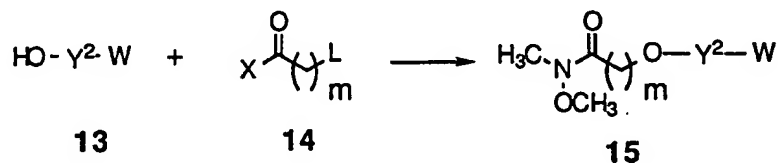


In Step 4, compound **11** is reacted with an amine of formula **6** in a manner such as that described in Procedure A1, Step 2.

Procedure C:

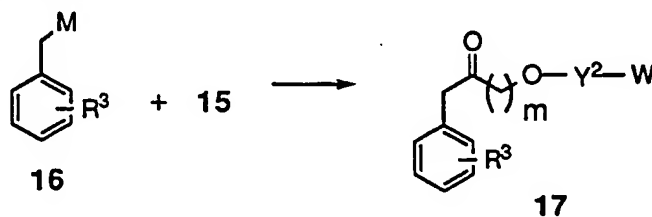
- 5 Compounds of formula I wherein $-Q-(C(R^7)(R^8))_s-$ is $-N(R^{11})(CH_2)_s-$ and s is 2 or 3, Z^1 is hydrogen, Y^1 is $-C(=NO(C(R^9)(R^{10}))_rT)(CH_2)_m-$ and m is 1 to 3, and J is $-O-$ can be prepared as shown in the following reaction scheme:

Step 1



- 10 In step 1, an alcohol with the structure **13** is treated with a base such as NaH or LDA, preferably NaH, in an inert solvent such as THF or DMF. The resulting alkoxide is added to an electrophile such as a compound of formula **14** wherein m is 1, 2 or 3 and L is a leaving group as defined above, preferably Br, and X is $-N(CH_3)OCH_3$ or $-Oalkyl$.
- 15 Preferable reaction temperatures range from -20 to 50°C . When X is $Oalkyl$, it is then treated with $HN(CH_3)OCH_3$ and $Al(CH_3)_3$ in an inert organic solvent such as THF or toluene at a temperature of -20 to 40°C .

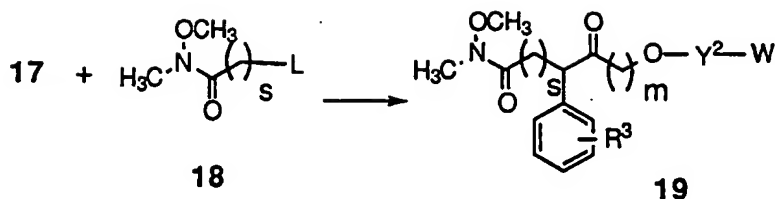
Step 2



- 20 In step 2, compound **15** is treated with a reagent of formula **16** wherein M is Li, MgCl or MgBr, in an inert organic solvent such as THF or ether, at a temperature of -78 to 40°C .

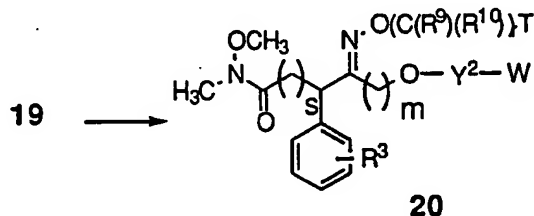
- 14 -

Step 3



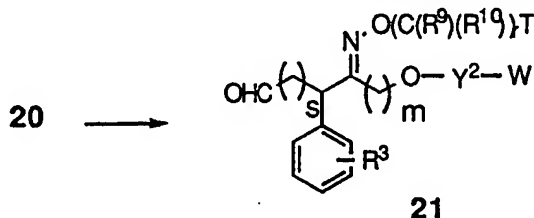
In step 3, compound 17 is treated with a suitable base such as $\text{NaN}(\text{TMS})_2$ or LDA, preferably $\text{NaN}(\text{TMS})_2$, in an inert organic solvent such as THF. The resulting anion is treated with a compound of formula 18 wherein s is 1 or 2 and L is a leaving group as defined above, in an inert organic solvent such as THF or ether at reaction temperatures ranging from -78 to 30°C .

Step 4



A compound of formula 19 is converted to the corresponding oxime of formula 20 by treating the ketone 19 with a hydroxylamine derivative of the formula $\text{H}_2\text{NO}(\text{C}(\text{R}^9)(\text{R}^{10}))_r\text{T}$, in a suitable organic solvent such as pyridine at a temperature of from 25 to 100°C . Alternatively, a low molecular weight alcohol (e.g. CH_3OH or $\text{CH}_3\text{CH}_2\text{OH}$) can be used as the solvent, in which case a base such as sodium acetate must be added.

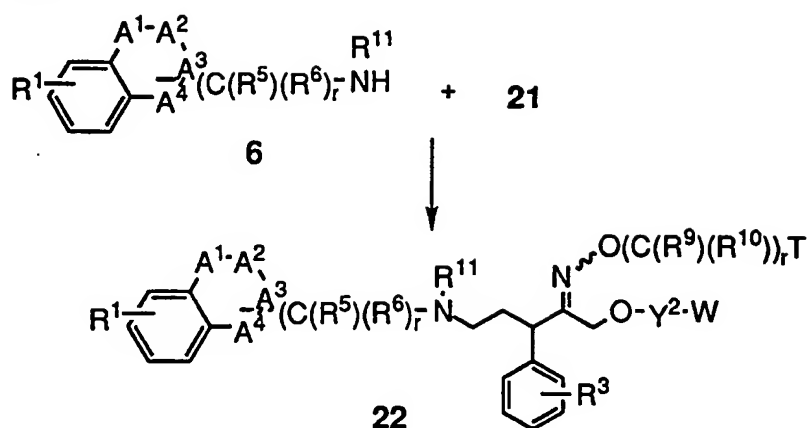
Step 5



In step 5, compound 20 is treated with a reducing agent such as DIBAL-H in an inert organic solvent such as THF or CH_2Cl_2 at a temperature from -78 to -40°C .

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Step 6

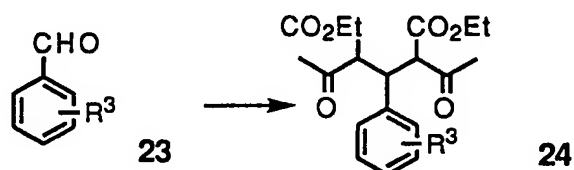


In Step 6, compound **21** is reacted with an amine of formula **6** in a manner such as that described in Procedure A1, Step 2.

Procedure D:

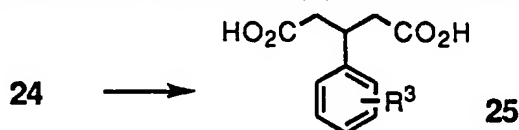
- 5 Compounds of formula I wherein $-\text{Q-(C(R}^7\text{)(R}^8\text{))}_s-$ is $-\text{N(R}^{11}\text{)CH}_2\text{CH}_2-$, Z^1 is hydrogen, Y^1 is $-\text{CH}_2\text{C(O)-}$, and J is $-\text{N(CH}_3\text{)-}$ can be prepared as shown in the following reaction scheme:

Step 1



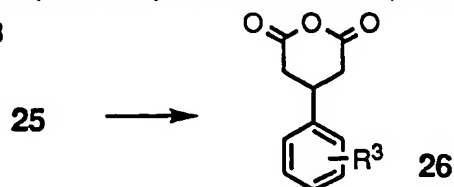
- 10 In step 1, an aldehyde of formula **23** is reacted with ethylacetoacetate in polar organic solvent such as $\text{CH}_3\text{CH}_2\text{OH}$ in the presence a suitable base, e.g., piperidine, at a temperature of 10 to 50 °C.

Step 2



- 15 In step 2, compound **24** is converted to the diacid compound of formula **25** by treatment with a strong base such as NaOH in an aqueous alcoholic solvent such as $\text{CH}_3\text{CH}_2\text{OH}$ at a temperature from 60 to 100°C, preferably at the reflux temperature of the solvent.

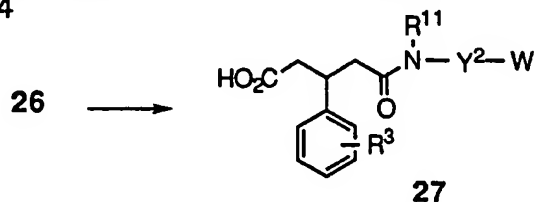
Step 3



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In step 3, compound **25** is treated with a dehydrating reagent such as CH_3COCl or DCC, preferably CH_3COCl .

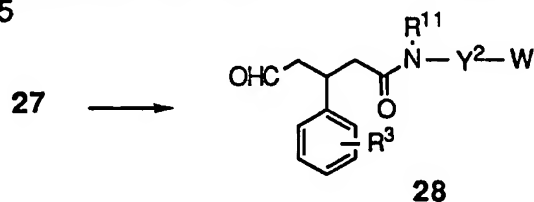
Step 4



Treatment of anhydride **26** with an amine of the formula $\text{W}-\text{Y}^2-\text{NHR}^{11}$

- 5 $\text{Y}^2-\text{NHR}^{11}$ in a suitable solvent such as CH_2Cl_2 in the presence of a suitable base such as Et_3N or N,N -dimethylamino-pyridine gives acid **27**.

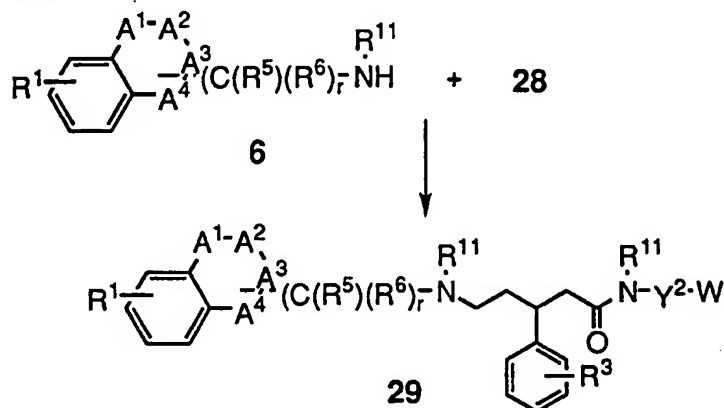
Step 5



Acid **27** is converted to the alcohol by a suitable reduction procedure. For example, a compound **27** is treated with 1,1'-carbonyl-diimidazole in an organic solvent such as ethyl acetate in the presence of a suitable base such as N,N -dimethylaminopyridine followed by treatment with aqueous NaBH_4 . The alcohol is converted to the aldehyde **28** by a suitable oxidation procedure.

10

Step 6

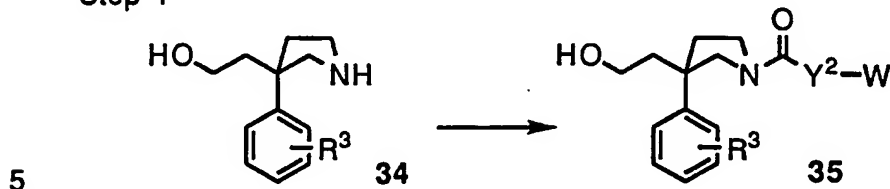


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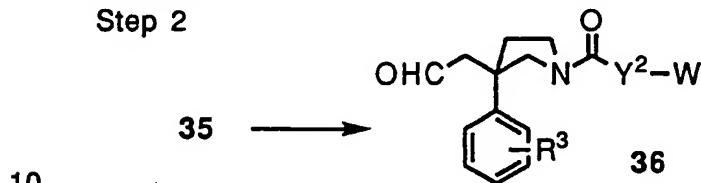
In Step 6, compound **28** is reacted with an amine of formula **6** in a manner such as that described in Procedure A1, Step 2.

Procedure E:

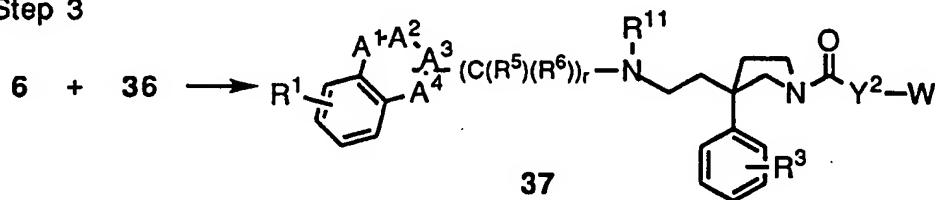
Compounds of formula I wherein $-Q-(C(R^7)(R^8))_s-$ is $-N(R^{11})CH_2CH_2-$, Y^1 is $-CH_2-$, J is $-N(Z^2)CO-$ and Z^1 and Z^2 together are $-CH_2CH_2-$ can be prepared as shown in the following reaction scheme:

Step 1

In step 1, 3-(R^3 -phenyl)-3-(2-hydroxyethyl)-pyrrolidine **34** (which can be obtained by the procedure described in T.B. Burkholder et al. *Bioorg. & Med. Chem. Let.* 6, (1996), p. 951) is acylated by standard procedures as described in Procedure A2, Step 4.

Step 2

In step 2, compound **35** is converted to aldehyde **36** by a suitable oxidation procedure, for example by the Swern procedure.

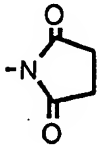
Step 3

In step 3, compound **36** is reacted with an amine of formula **6** in a procedure such as described in Procedure A1, Step 2.

Reactive groups not involved in the above processes can be protected during the reactions with conventional protecting groups which can be removed by standard procedures after the reaction. The following Table 1 shows some typical protecting groups:

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Table 1

Group to be Protected	Group to be Protected and Protecting Group
-COOH	-COOalkyl, -COObenzyl, -COOphenyl
>NH	$\begin{array}{l} \text{>NCOalkyl, >NCObenzyl, >NCOphenyl,} \\ \text{>NCH}_2\text{OCH}_2\text{CH}_2\text{Si(CH}_3\text{)}_3, \text{ >NC(O)OC(CH}_3\text{)}_3, \\ \text{>N-benzyl, >NSi(CH}_3\text{)}_3, \text{ >NSi(CH}_3\text{)}_2\text{C(CH}_3\text{)}_2\text{CH}_3 \end{array}$
-NH ₂	
-OH	$\begin{array}{l} \text{-OCH}_3, \text{-OCH}_2\text{OCH}_3, \text{-OSi(CH}_3\text{)}_3, \text{-OSi(CH}_3\text{)}_2\text{C(CH}_3\text{)}_2\text{CH}_3 \\ \text{or -OCH}_2\text{phenyl} \end{array}$

Compounds of formula I have been found to be antagonists of NK₁ and/or NK₂ and/or NK₃ receptors, and are therefore useful in treating conditions caused or aggravated by the activity of said receptors.

The present invention also relates to a pharmaceutical composition comprising a compound of formula I and a pharmaceutically acceptable carrier. Compounds of this invention can be administered in conventional oral dosage forms such as capsules, tablets, powders, cachets, suspensions or solutions, or in injectable dosage forms such as solutions, suspensions, or powders for reconstitution. The pharmaceutical compositions can be prepared with conventional excipients and additives, using well known pharmaceutical formulation techniques. Pharmaceutically acceptable excipients and additives include non-toxic and chemically compatible fillers, binders, disintegrants, buffers, preservatives, anti-oxidants, lubricants, flavorings, thickeners, coloring agents, emulsifiers and the like.

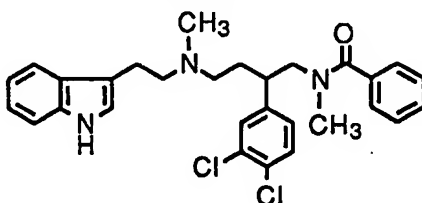
The daily dose of a compound of formula I for treating asthma, cough, bronchospasm, inflammatory diseases, migraine, nociception and gastrointestinal disorders is about 0.1 mg to about 20 mg/kg of body weight per day, preferably about 0.5 to about 15 mg/kg. For an average body weight of 70 kg, the dosage range is therefore from about 1 to about 1500 mg of drug per day, preferably about 50 to about

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200 mg, more preferably about 50 to about 500 mg/kg per day, given in a single dose or 2-4 divided doses. The exact dose, however, is determined by the attending clinician and is dependent on the potency of the compound administered, the age, weight, condition and response of the patient.

Following are examples of preparing starting materials and compounds of formula I.

Example 1



10

Step 1: Reflux a solution of 3-(3,4-dichlorophenyl)-2-propenoic acid (26.02 g, 0.12 mol), CH₃OH (500 mL) and saturated methanolic HCl (5.0 mL) for 14 h. Cool to precipitate white crystals and collect by filtration to give 22.6 g of the methyl ester product (0.098 mol, 82%). Concentrate the filtrate and purify on a short silica gel column, eluting with 3:1 hexane: EtOAc to obtain an additional 4.43 g of the methyl ester (0.019 mol, 16%).

Step 2: Treat the product of Step 1 (25.0 g, 0.108 mol) with 40% methanolic benzyltrimethylammonium hydroxide (9.3 mL, 22 mmol) and CH₃NO₂ (210 mL) and heat at 80°C for 1.5 h. Dilute the reaction mixture with diethyl ether (Et₂O) (1 L), wash with 1N HCl (400 mL and 100 mL) and brine (500 mL), dry with MgSO₄ and concentrate. Chromatograph on silica gel, eluting with 3:1 hexane:EtOAc, to obtain 31.0 g of methyl 3-(3,4-dichlorophenyl)-4-nitrobutyrate (0.106 mol, 98%).

Step 3: To a 1M Et₂O solution of LiAlH₄ (200 mL, 200 mmol) at 0°C, slowly add the product of Step 2 (14.35 g, 49.1 mmol) dissolved in tetrahydrofuran (THF) (100 mL). Allow the reaction mixture to warm to room temperature and stir for 45 min. After recooling to 0°C, quench the excess LiAlH₄ by the careful addition of aqueous saturated Na₂SO₄ (20 mL). Dry the solution with Na₂SO₄ and filter. Wash the lithium salts with Et₂O (3 x 300 mL). Concentrate the combined filtrates to give 8.65 g of 4-amino-3-(3,4-dichlorophenyl)-butanol as a white solid (36.9 mmol, 75%).

Step 4: Treat a solution of the amino alcohol from Step 3 (8.13 g, 34.7 mmol) and imidazole (3.56 g, 52.3 mmol) in CH₂Cl₂ (350 mL) with

- 20 -

5 *tert*-butyldimethylsilyl chloride (7.84 g, 52.0 mmol). After stirring for 1 h, wash the reaction with 0.1 N HCl (350 mL) and extract the aqueous layer with CH₂Cl₂ (100 mL). Wash the combined organic layers with saturated NaHCO₃ (200 mL) and brine (200 mL), dry with K₂CO₃, and concentrate to give 11.4 g of the silylated product as a pale yellow viscous liquid (34.2 mmol, 98 %).

10 Step 5: Treat the amine from step 4 (15.3 g, 44 mmol) in CH₂Cl₂ (250 mL) with triethyl amine (Et₃N) (13 mL, 93 mmol) and benzoyl chloride (8.0 mL, 69 mmol). Stir at room temperature for 1.5 h, then wash the reaction mixture with H₂O (500 mL) and extract the aqueous layer with CH₂Cl₂ (2 x 100 mL). Wash the combined organic layers with 0.3 N HCl (150 mL), saturated NaHCO₃ (150 mL) and brine (200 mL), dry with MgSO₄ and concentrate. Chromatograph the crude material on silica gel, eluting with 6:1 to 3:1 hexane:EtOAc to give 13.7 g of the benzamide (30 mmol, 69%).

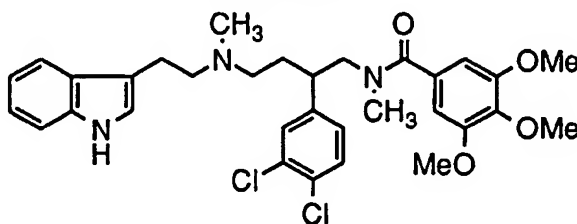
20 Step 6: Treat the product of step 5 (10.54 g, 23.3 mmol) in THF (170 mL) with 60 % NaH (1.87 g, 47 mmol) followed by CH₃I (1.9 mL, 30.5 mmol). Heat the reaction mixture at 60°C for 30 min and then partition between Et₂O (250 mL) and H₂O (500 mL). Extract the aqueous layer with Et₂O (500 mL), wash the combined organic layers with brine (250 mL), dry with MgSO₄ and concentrate to give 9.9 g of the methyl benzamide product as a colorless oil (21 mmol, 91%).

25 Step 7: Treat the product of Step 6 (9.9 g, 21 mmol) in THF (125 mL) with a 1 M THF solution of *t*-butylammonium fluoride (50 mL, 50 mmol) and stir for 4 h. Partition the reaction mixture between H₂O (200 mL) and Et₂O (100 mL). Extract the aqueous layer with Et₂O (2 x 100 mL), combine the organic layers and wash with brine (100 mL), dry with MgSO₄ and concentrate. Purify the crude product by recrystallization in CH₂Cl₂:hexane to give 5.5 g of the alcohol product as white crystals (15.6 mmol, 74.4 %).

30 Step 8: Stir the product of step 7 (1.37 g, 3.8 mmol) in CH₂Cl₂ (20 mL) with Et₃N (0.6 mL, 4.3 mmol) and C H₃SO₂Cl (0.32 mL, 4.1 mmol) for 45 min, then add the reaction mixture to 0.3 N HCl (200 mL) and extract with CH₂Cl₂ (3 x 100 mL). Wash the combined organic layers with saturated NaHCO₃ (100 mL) and brine (100 mL), dry with MgSO₄ and concentrate to obtain 1.49 g of the methanesulfonate ester as a colorless viscous oil (3.47 mmol, 89%).

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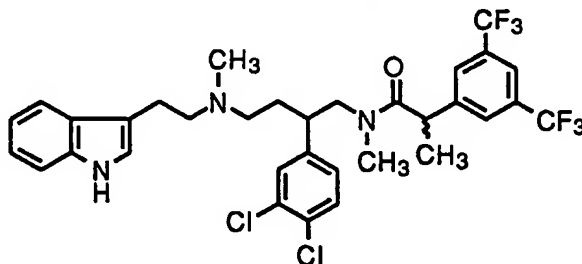
Step 9: Heat the product of step 8 (0.499 g, 1.16 mmol) in DMF (5 mL) with Nω-methyl tryptamine (0.241 g, 1.38 mmol) and NaI (16.4 mg, 0.11 mmol) at 50°C for 38 h. Add the reaction mixture to saturated NaHCO₃ (100 mL) and extract with CH₂Cl₂ (3 x 40 mL), wash the combined organic layers with brine (80 mL), dry with MgSO₄ and concentrate. Chromatograph on silica gel, eluting with 20:1:0.1 to 13:1:0.1 CH₂Cl₂:MeOH:NH₃ (aq.), gave 413 mg of the title compound as a white foam (0.81 mmol, 70 %). HRMS (FAB, M+H⁺): m/e calc'd for [C₂₉H₃₂Cl₂N₃O]⁺ 508.1922; found 508.1929.

Example 2

Step 1: To a 0°C solution of the product of Example 1, Step 4 (8.0 g, 22.9 mmol) in DMF, add 4-methylmorpholine (NMM) (2.5 mL, 22.9 mmol), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (6.5 g, 34.4 mmol), 1-hydroxybenzotriazole hydrate (HOBT) (3.0 g, 22.9 mmol) and 3,4,5-trimethoxybenzoic acid (4.9 g, 22.9 mmol). Stir the reaction mixture at 0°C for 30 min., then at room temperature overnight. Concentrate under high vacuum and resuspend the resulting material in H₂O and extract with EtOAc. Wash the combined organic layers with saturated NaHCO₃, dry with MgSO₄ and concentrate. Chromatograph the crude product on silica gel, eluting with 20:1 CH₂Cl₂:NH₃ saturated CH₃OH to give 9.0 g of amide as a light yellow solid (72 %).

Steps 2-5: Convert the product of Step 1 to the title compound using procedures similar to those described in Example 1, Steps 6 to 9.

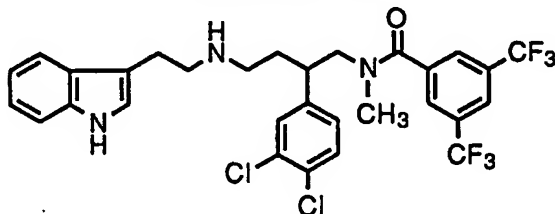
MS (FAB, M+H⁺): m/e 598; **Analysis for C₃₂H₃₇Cl₂N₃O₄:** Calc'd C, 64.21; H, 6.23; N, 7.20%; Found: C, 63.84; H, 6.37; N, 7.00.

Example 2a

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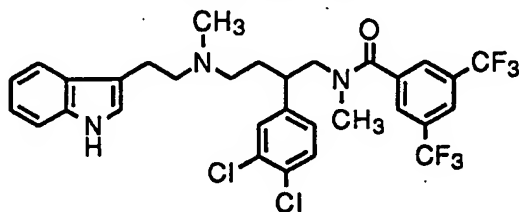
Using the procedure of Example 2, substitute 3,5-bis(trifluoromethyl)phenylacetic acid for 3,4,5-trimethoxybenzoic acid in step 1, and in step 2, methylate the benzylic position of the phenyl acetyl group and the amide position to obtain the title compound. MS (FAB, M+H⁺): m/e 672.

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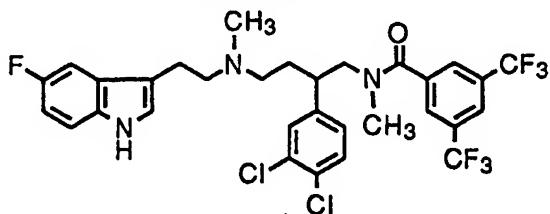
Example 2b

Using the procedure of Example 2, substitute 3,5-bis(trifluoromethyl)benzoic acid for 3,4,5-trimethoxybenzoic acid in step 1, and tryptamine for Nω-methyltryptamine in step 5 to obtain the title compound.

10 HRMS (FAB, M+H⁺): m/e calc'd for [C₃₀H₂₈N₃OC₂F₆]⁺ 630.1514; found 630.1513.

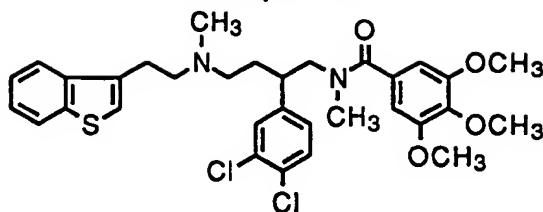
Example 2c

15 Using the procedure of Example 2, substitute 3,5-bis(trifluoromethyl)benzoic acid for 3,4,5-trimethoxybenzoic acid in step 1 to obtain the title compound.

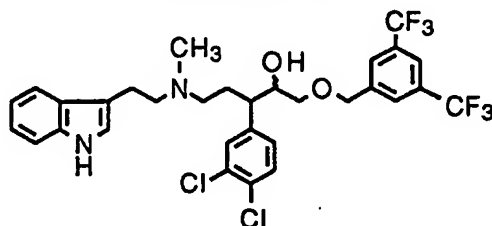
Example 2d

20 Using the procedure of Example 2c, substitute 5-fluoro-Nω-methyltryptamine for Nω-methyltryptamine in step 5 to obtain the title compound. HRMS (FAB, M+H⁺): m/e calc'd for [C₃₁H₂₉N₃OC₂F₇]⁺ 662.1576; found 662.1584.

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Example 2e

- Using the procedure of Example 2, substitute N-methyl-3-(2-aminoethyl)benzothiophene for N-methyltryptamine in step 5 to obtain the title compound. HRMS (FAB, M+H⁺): m/e calc'd for [C₃₂H₃₇N₂O₄Cl₂S]⁺ 615.1851; found 615.1850.

Example 3

- Step 1:** Cool a solution of 3-(3,4-dichlorophenyl)-2-propenoic acid (100 g, 461 mmol) in dry DMF (500 mL) to 0 °C and treat with Cs₂CO₃ (100 g, 307 mmol). Stir the resulting off-white slurry for 15 min., then add CH₃I (33 mL, 530 mmol) via syringe. After 1 h, add additional DMF (250 mL) and stir the slurry for 14 h, then partition between ethyl acetate (EtOAc) (1.5 L) and half saturated aqueous NaHCO₃ (500 mL). Separate the organic layer and extract the aqueous layer with EtOAc (1 L, 500 mL). Wash the combined organic layers with half saturated aqueous NaHCO₃ (500 mL) and water (5 x 500 mL), then dry over Na₂SO₄ and concentrate to obtain methyl 3-(3,4-dichlorophenyl)-2-propenoate, 105.4 g (456 mmol, 99%), as light brown needles.
- Step 2:** Treat a solution of the product of Step 1 (15 g, 65 mmol) in dry THF (250 mL) (kept cool in a large ambient temperature water bath) with DIBAL-H (140 mL, 140 mmol) over 30 min. Stir the resulting solution for 30 min at 23 °C, pour into Et₂O (500 mL) and treat with water (5 mL), 15 % NaOH (5 mL) and water (15 mL). Stir for 5 min, dilute the mixture with Et₂O (200 mL), add 15 % NaOH (15 mL), then add MgSO₄ to obtain a colorless precipitate. Remove the aluminum salts by filtration through a coarse glass frit, wash the solids with Et₂O (1 L) and concentrate the filtrate *in vacuo* to give 3-(3,4-dichlorophenyl)-2-propene-1-ol as an off-white solid, 13.2 g (65 mmol, 99%).

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Step 3: Combine a solution of the product of Step 2 (13.2 g, 65 mmol) in CH_2Cl_2 (250 mL) at 0°C with pyridine (7.89 mL, 97.5 mmol) and dimethylaminopyridine (DMAP) (397 mg, 3.25 mmol), followed by acetyl chloride (6.48 mL, 74.7 mmol). Allow the mixture to warm to 23°C , pour
5 into 1 M HCl (100 mL) and wash the resulting organic layer with 1 M HCl (100 mL) followed by water (5 x 100 mL; pH=6.5-7). Dry the organic layer over Na_2SO_4 and concentrate, providing 3-(3,4-dichloro-phenyl)-2-propene-1-ol acetate as a colorless oil, 15.4 g (62.9 mmol, 97%).

Step 4: Treat a solution of the product of Step 3 (15 g, 61 mmol,
10 dried by azeotropic distillation with toluene, 1 x 50 mL) in dry THF (250 mL) at -78°C with chlorotriethylsilane (20.2 mL, 120 mmol), rapidly followed by the addition of 0.5 M toluene solution of potassium bis(trimethylsilyl)amide (183 mL, 91.5 mmol) via addition funnel over 50 min. Allow the mixture to warm to 23°C , then heat to reflux for 3 h. Allow the
15 solution to gradually cool overnight, then quench with saturated NH_4Cl (150 mL). Stir the resulting mixture vigorously for 3h, treat with 1M HCl (150 mL) and extract with Et_2O (500 mL). Extract the aqueous layer with Et_2O (400 mL) and wash the combined organic layers with 300 mL of 5% NaOH followed by 8 x 150 mL of 5 % NaOH. Cool the combined aqueous
20 layers to 5°C and carefully (temperature kept to $5-10^\circ\text{C}$) acidify with conc. HCl (ca 175 mL) to pH 1. Extract the aqueous layer with CH_2Cl_2 (2 x 800 mL), dry (Na_2SO_4) and concentrate to give 3-(3,4-di-chlorophenyl)-4-pentenoic acid as a faint yellow oil, 13.4 g (54.5 mmol, 89%).

Step 5: Treat a solution of the product of Step 4 (5.0 g, 20.4 mmol) in
25 dry CH_2Cl_2 (60 mL) with purified *m*-CPBA (7 g, 40 mmol) [wash 13 g of commercial 55% *m*CPBA in 250 mL of benzene with pH 7.4 buffer (5 x 30 mL), dry (Na_2SO_4) and concentrate to give about 9 g of pure *m*-CPBA]. After stirring for 48 h, add Amberlyst 15 (1.2 g) and stir the mixture for 8 h. Remove the Amberlyst by filtration through a medium porosity glass frit,
30 rinsing with EtOAc. Wash the filtrate with 100 mL of saturated Na_2SO_3 / NaHCO_3 (1:1), dry the resulting organic layer and concentrate *in vacuo*. Take up the crude product in hexane: CH_2Cl_2 (1:1) and filter to give 3.3 g (12.6 mmol, 62%) of a mixture of isomers (3:2, *trans* / *cis*) of the title compound as a colorless soft solid. Concentrate the filtrate to give 2.0 g of
35 a viscous oil which is purified by silica gel chromatography (column: 7 x 15 cm; solvent: hexane:EtOAc, 5:4 gradient to 1:1) to give 1.07 g (4.1 mmol, 20%) of the pure *cis* isomer of the title compound as an oil, total yield 4.3 g (16.47 mmol, 81%).

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Step 6: Treat a solution of 4-(3,4-dichlorophenyl)-dihydro-5-(hydroxymethyl) 2(3H)-furanone (3.3 g, 12.6 mmol, 3:2 ratio of stereo-isomers by NMR) in dry DMF (10 mL) with 3,5-bis(trifluoromethyl)benzyl bromide (5.9 mL, 32.2 mmol) followed by Ag₂O (5.8 g, 25.3 mmol); wrap the vessel in foil and stir for 2.5 d. Apply the crude material to a pad of silica gel (10 cm x 4 cm) packed with hexane:EtOAc (1:1), washing the pad with the same solvent until no further product is eluted as shown by TLC; concentrate the resulting filtrate *in vacuo* to give the crude product as a solid (10g). Dissolve the crude was dissolved in hexane:EtOAc (4:1) and purify by silica gel chromatography (column: 7.5 x 19; solvent: hexane:EtOAc, 4:1) to give 3.33 g (6.8 mmol, 54%) of isomer (*trans*)-[[[(3,5-bis(trifluoromethyl)phenyl)methoxy)methyl]-4-(3,4-dichlorophenyl)-dihydro-2(3H)-furanone and 1.08 g (2.2 mmol, 17%) of the corresponding cis isomer for a total yield of 71%.

Trans isomer: HRMS (FAB, M+H⁺): *m/e* calc'd for [C₂₀H₁₅O₃Cl₂F₆]⁺: 487.0302, found 487.0312.

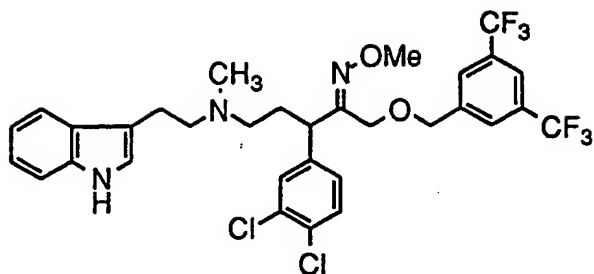
Cis isomer: HRMS (FAB, M+H⁺): *m/e* calc'd for [C₂₀H₁₅Cl₂F₆O₃]⁺: 487.0302, found 487.0297.

Step 7: Cool a solution of the cis isomer of Step 6 (2.1g, 4.31 mmol) in dry CH₂Cl₂ (50 mL) to -78 °C and treat with DIBAL-H (5.1 mL, 5.1 mmol; 1M in CH₂Cl₂). Stir for 2 h at -78 °C, then treat the solution with NaF (905 mg 22 mmol) and water (400 µL, 22 mmol, 5 eq). Allow the suspension to warm to 23 °C and stir for 45 min., dilute the mixture with ET₂O (50 mL) and filter through a pad of silica gel (6.5 cm x 2 cm; 150 mL vacuum glass frit) packed with hexane:EtOAc (1:1)), washing the pad with hexane:EtOAc (1:1) until no further product is evident as shown by TLC. Concentrate the filtrate to give 1.92 g (3.86 mmol, 91%) of the title compound as a foam which was used without further purification in the next step.

Step 8: Stir a suspension of the lactol of Step 7 (1.03 g, 2.1 mmol), Nω-methyl tryptamine (0.73 g, 4.2 mmol) and 3A molecular sieves (1.7 g) in CF₃CH₂OH (5.0 mL) for 1 h, add NaCNBH₃ (0.26 g, 4.2 mmol) and stir the reaction mixture over the weekend, then concentrate. Chromatograph the residue on silica gel, eluting with 20:1 CH₂Cl₂/CH₃OH (saturated with ammonia) to give 0.809 g of the desired compound as a white solid (60 %). MS (FAB, M+H⁺): *m/e* 647.

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Example 4



- Step 1:** Combine 3,5-bis(trifluoromethyl)benzyl alcohol (25.0 g, 0.103 mol) in THF (50 mL) with 60% NaH (4.14 g, 0.104 mol) and stir for 30 min. Transfer the resulting alkoxide mixture via cannula over 30 min to a solution of methylbromoacetate (11.8 mL, 0.125 mol) in THF (250 mL). Stir the reaction mixture for 18h and add to 0.3 N HCl (300 mL). Separate the organic layer and extract the aqueous layer with EtOAc (2 x 150 mL). Wash the combined organic layers with brine (200 mL), dry with MgSO₄ and concentrate. Chromatograph on silica gel, eluting with 10:1 hexane: EtOAc, to give the desired product as a colorless liquid (26.1 g, 80%).
- Step 2:** Combine a suspension of N,O-dimethylhydroxylamine hydrochloride (19.5 g, 0.20 mol) in THF (350 mL) at 0°C with AlMe₃ (100 mL, 0.20 mol, 2M in toluene) and allow to warm to room temperature for 30 min. Cool this mixture to 0°C, add the product of Step 1 (26.1 g, 82.7 mmol) dissolved in THF (140 mL) over 30 min., warm the mixture to room temperature and stir for 40 min. Cool the reaction mixture to 0°C, quench by the careful addition of 1N HCl (100 mL) and concentrate. Partition the residue between H₂O (250 mL) and CH₂Cl₂ (300 mL), extract the aqueous layer with CH₂Cl₂ (2 x 150 mL) dry the combined organic layers with MgSO₄ and concentrate. Chromatograph on silica gel, eluting with 3:1 hexane:EtOAc to give the product as a colorless liquid (28.1 g, 98%).
- Step 3:** Treat a suspension of Mg (1.35 g, 55.6 mmol) in Et₂O (10 mL) maintained at 30°C with a solution of α,3,4-trichlorotoluene (7.7 mL, 55.6 mmol) in Et₂O (45 mL) over 1h and then stir for 30 min. Transfer the resulting solution via cannula over 30 min to a -78°C solution of the product of Step 2 (9.56 g, 27.7 mmol) in Et₂O (350 mL) and warm the reaction mixture to room temperature over 1 h. Cool the reaction mixture to 0°C and treat with HCl saturated CH₃OH (4 mL) and CH₃OH (100 mL), stir for 1 h and concentrate. Partition the residue between CH₂Cl₂ (300

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mL), Et₂O (500 mL) and H₂O (250 mL), dry the organic layer with MgSO₄ and concentrate. Chromatograph on silica gel, eluting with 2:1 CH₂Cl₂:hexane, to give the product as a white solid (6.22 g, 50%).

Step 4: To a solution of sodium bis(trimethylsilyl)amide (2.5 mmol) in THF (20 mL) at -78°C, add, dropwise, a solution of the product from Step 3 (1.11 g, 2.5 mmol) in THF (5 mL) over 15 min. Stir at -78°C for 2 h and then add a solution of 2-bromo-N-methoxy-N-methylacetamide (455 mg, 2.5 mmol) in THF (5 mL) dropwise over 10 min. Allow the mixture to warm to room temperature over 1 h, then stir at room temperature for an additional 30 min. Add brine to the reaction mixture (2 mL) and concentrate. Partition the residue between CH₂Cl₂ (60 mL), Et₂O (90 mL) and H₂O (30 mL). Wash the organic layer with brine (30 mL), dry with Na₂SO₄ and concentrate. Chromatograph on silica gel, eluting with CH₂Cl₂ to give the product as a colorless oil (800 mg, 56%).

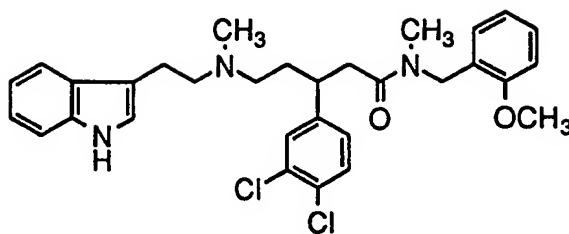
Step 5: Heat a mixture of the product of Step 4 (597 mg, 1.1 mmol), pyridine (10 mL) and methoxylamine hydrochloride (101.5 mg, 1.2 mmol) at 60°C for 1 h and then concentrate. Chromatograph on silica gel, eluting with 3:1 hexane:EtOAc to obtain the oxime (syn-isomer) as a colorless oil (442 mg, 70%). HRMS (FAB, M+H⁺): *m/e* calc'd for [C₂₃H₂₂Cl₂F₆N₂O₄]⁺: 575.0939, found 575.0932.

Step 6: Treat the product of Step 5 (347 mg, 0.60 mmol) in THF (6 mL) at -78°C with DIBAL-H (1.8 mL, 1.8 mmol, 1M in hexane) over 15 min. Add H₂O (5 mL) and warm to room temperature; to this mixture, add NAF (0.2 g, 4.8 mmol) and stir for 20 min. Partition the reaction mixture between brine (25 mL) and Et₂O (25 mL), dry the organic layer with MgSO₄ and concentrate to give the crude product (298 mg, 94%).

Step 7: Treat the aldehyde of Step 6 (35 mg, 68 µmol) with Nω-methyl tryptamine (26.2 mg, 0.15 mmol), CF₃CH₂OH (0.4 mL) and crushed 3Å molecular sieves (36 mg) and stir for 20 min. Add NaCNBH₃ (13 mg, 0.21 mol), stir for 14 h and concentrate. Chromatograph on silica gel, eluting with 10:1:0.1 CH₂Cl₂:MeOH:NH₃(aq) to give the title compound as a colorless oil (32 mg, 70%). MS (FAB, M+H⁺): *m/e* calc'd 674.1776; found 674.1787.

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Example 5



- Step 1:** Combine 3,4-dichlorobenzaldehyde (100 g) in 95% $\text{CH}_3\text{CH}_2\text{OH}$ (120 mL) with ethylacetoacetate (146 mL) and stir until a homogenous solution is obtained. Treat this solution with piperidine (8 mL) and allow to stand for 18 hours. Recrystallize the crude product from 95% ethanol to give diethyl-3,4-dichlorobenzal-*bis*-acetoacetate (230 g).
- Step 2:** Reflux the product of Step 1 (155 g) in $\text{CH}_3\text{CH}_2\text{OH}$ (2 L) and 50% NaOH (2 l) for 4 hours. Add water (1 L) to the reaction mixture and remove approx. 1.5 L of solvent by distillation. Pour the remaining solution onto ice (1 Kg) and add sufficient HCl to adjust the pH to 1. Extract the resulting solution with EtOAc (3 X 1.5 L), dry the combined extracts over MgSO_4 , filter and concentrate to give 100 g of 3-(3,4-dichlorophenyl)-glutaric acid.
- Step 3:** Heat a combination of the product of Step 2 (100 g) and acetyl chloride (300 mL) at reflux for 5 hours. Cool the reaction mixture, azeotrope with toluene, and concentrate under reduced pressure. Slurry the residue with Et_2O (250 mL) and filter to afford 3-(3,4-dichlorophenyl)-glutaric anhydride (86 g).
- Step 4:** Sequentially treat the product of Step 3 (5.9 g) in CH_2Cl_2 (80 mL) at 0°C with N-methyl-N-[2-(methoxyphenyl)methyl]amine (3.8 g), Et_3N (3.5 mL) and DMAP (278 mg). Stir the mixture at 0°C for 2 h, allow to warm to room temperature and stir for 20 h. Wash the reaction mixture with 1N HCl (1 x 100 mL) and brine (1 x 100 mL), dry the organic layers over MgSO_4 , filter and concentrate to afford 3,4-dichloro- β -[2-[[2-(methoxyphenyl)methyl]-methylamino]-2-oxoethyl] benzenepropanoic acid (9.3 g).
- Step 5:** Treat the acid from Step 4 (9.3 g) in EtOAc (100 mL) with 1,1'-carbonyldiimidazole (4.62 g) and DMAP (345 mg), stir the resulting solution at room temperature for 15 min., then heat at 50°C for 1 h. Cool the reaction mixture to 0°C and treat with a solution of NaBH_4 (3.45 g) in H_2O (50 mL), warm slowly to room temperature and stir for 12 h. Dilute the reaction mixture with EtOAc (250 mL) and wash with 1N HCl (1 x 100

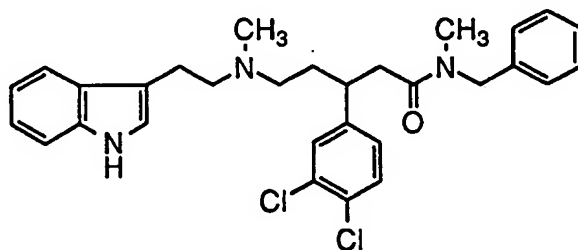
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- mL) and H₂O (1 x 100 mL), dry over MgSO₄, filter and concentrate under reduced pressure to yield a crude oil (13 g). Chromatograph on silica gel, eluting with 5% CH₃OH/CH₂Cl₂ to give 3,4-dichloro-β-(2-hydroxyethyl)-N-methyl-N-[(2-methoxyphenyl) methyl]benzenepropanamide (8.7 g).
- 5 HRMS (FAB, M+H⁺): *m/e* cal'd for [C₂₀H₂₄NO₃Cl₂]⁺ 396.1133; found 396.1124.

- Step 6:** Cool a solution of oxalyl chloride (1.43 mL) in CH₂Cl₂ (30 mL) to -78°C, add DMSO (2.32 mL) dropwise over 15 mins., stir for 15 min., then add a CH₂Cl₂ (20 mL) solution of the product of Step 5 (1.3 g) over 20 min. Stir the mixture for 30 min, treat with Et₃N (9.2 mL) and stir for an additional 30 min. at -78°C, followed by 1.5 h at room temperature. Quench the reaction mixture with water and dilute with CH₂Cl₂ (100 mL). Separate the organic fraction, wash sequentially with 1N HCl (1 x 50 mL), sat. NaHCO₃ (1 x 50 mL) and brine (1 x 50 mL), dry over MgSO₄, filter and concentrate under reduced pressure to yield an oil. Chromatograph on silica gel, eluting with 50-100% EtOAc/hexane to give 3,4-dichloro-β-(2-oxoethyl)-N-methyl-N-[(2-methoxyphenyl)methyl] benzenepropanamide (950 mg).
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- Step 7:** Sequentially treat the aldehyde of Step 6 (770 mg), in 2,2,2-trifluoroethanol (5 mL) with molecular sieves 3A (510 mg), N-ω-methyltryptamine (510 mg) and NaCNBH₃ (123 mg). Stir the resulting mixture at room temperature for 18 h., dilute the reaction mixture with EtOAc (50 mL) and filter through silica gel. Concentrate the resulting organic layer under reduced pressure to give the crude product as an oil.
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25
- Chromatograph on silica gel, eluting with 10% MeOH/CH₂Cl₂ gave the title compound (525 mg). HRMS (FAB, M+H⁺): *m/e* cal'd for [C₂₁H₃₅N₃O₂Cl₂]⁺ 552.2185; found: 552.2179.

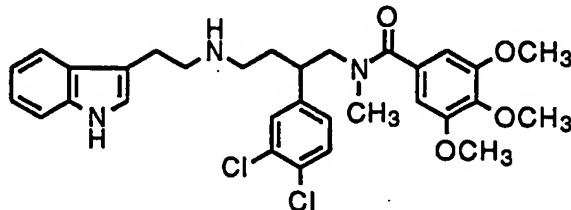
Example 6



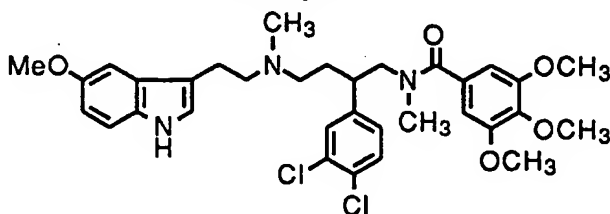
- 30 The compound of Example 6 is prepared by a procedure similar to that described in Example 5 except using N-methyl-N-phenyl-methylamine in place of N-methyl-N-[2-(methoxyphenyl)methyl]amine.

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MS (FAB, M+H⁺): *m/e* 522; Analysis for C₃₀H₃₃Cl₂N₃O·0.5H₂O: Found: C, 67.84; H, 6.43; N, 7.90; calc'd C, 67.79; H, 6.45; N, 7.91%.

Example 7

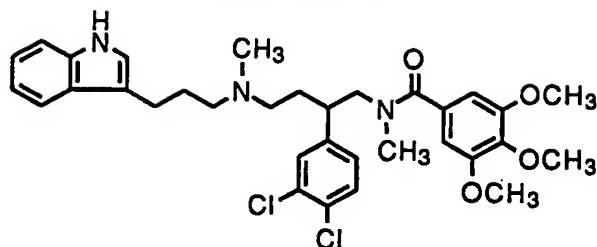
- 5 **Step 1:** To a -55°C solution of oxalyl chloride (2.25 mL, 25.8 mmol) in CH₂Cl₂ (45 mL), slowly add a solution of DMSO (2.4 mL, 33.8 mmol) in CH₂Cl₂ (45 mL) followed by a solution of the alcohol from Example 2, Step 3 (7.55 g, 17.1 mmol) in CH₂Cl₂ (90 mL). Stir for 30 min. at -55°C, add Et₃N (9.2 mL, 66 mmol) and stir for 2h at -55°C. Add 20% sat. KHSO₄
- 10 (75 mL) and Et₂O (100 mL), warm to room temperature and stir for 30 min. Add to Et₂O (225 mL) and remove the aqueous layer, wash with sat. NaHCO₃ (2 x 100 mL), H₂O (100 mL) and brine (100 mL). Dry the solution with MgSO₄ and concentrate to give 7.36 g of desired aldehyde product as a white foam (16.7 mmol, 98 %).
- 15 **Step 2:** Treat a solution of the product from step 1 (2.48 g, 5.6 mmol), tryptamine (1.44 g, 9.0 mmol) and crushed 3A mol. sieves (1.66 g) in CF₃CH₂OH (50 mL) with NaCNBH₃ (1.57 g, 25 mmol). Stir the reaction mixture for 30 min, filter and concentrate. Chromatograph the crude product on silica gel, eluting with 25:10:1 to 10:1:0.1 CH₂Cl₂:MeOH:NH₃
- 20 (aq.) to give 2.43 g of product as a white foam (4.16 mmol, 74 %).
MS (FAB, M+H⁺): *m/e* 584.

Example 7a

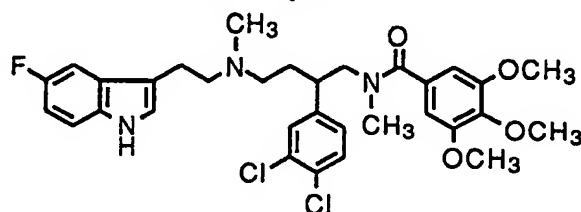
- 25 Using the procedure of Example 7, substitute 5-methoxy-N-ω-methyltryptamine for N-ω-methyltryptamine in step 2 to obtain the title compound.

HRMS (FAB, M+H⁺): *m/e* calc'd for [C₃₃H₄₀N₃O₅Cl₂]⁺ 628.2345; found 628.2345.

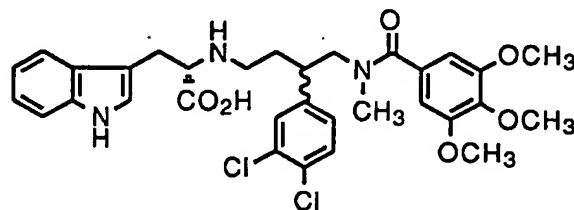
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Example 7b

- Using the procedure of Example 7, substitute 3-(N-methyl-3-aminopropyl)indole for N ω -methyltryptamine in step 2 to obtain the title compound.
- HRMS (FAB, M+H⁺): m/e calc'd for [C₃₃H₄₀N₃O₄Cl₂]⁺ 612.2396; found 612.2399.

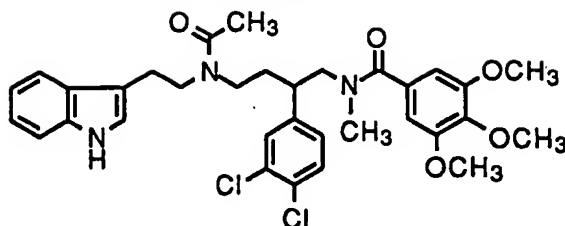
Example 7c

- Using the procedure of Example 7, substitute 5-fluoro-N ω -methyltryptamine for N ω -methyltryptamine in step 2 to obtain the title compound.
- HRMS (FAB, M+H⁺): m/e calc'd for [C₃₂H₃₇N₃O₄Cl₂F]⁺ 616.2145; found 616.2148.
- Separate the product of Example 7c into its enantiomers by chiral preparative HPLC using a Chiralpak AS column, eluting with 80:20 to 75:25 hexane:*i*-propylalcohol. The second enantiomer to elute has an optical rotation of $[\alpha]_D^{25} = +8.8^\circ$ (c= 4.4g/l in CH₃OH).

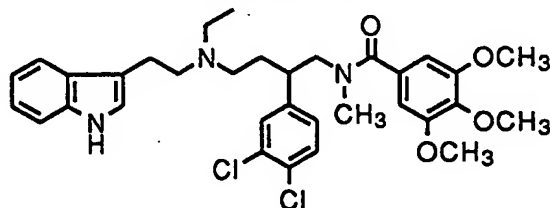
Example 7d

- Using the procedure of Example 7, substitute L-tryptophan for N ω -methyltryptamine in step 2 to obtain the title compound.
- MS (FAB, M+H⁺): m/e 628.

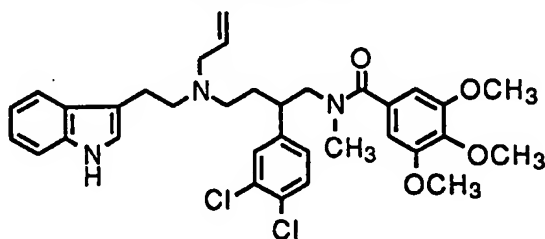
- 32 -

Example 8

- 5 Treat a solution of the product from Example 7 (144.8 mg, 0.248 mmol) and pyridine (30 μ L, 0.37 mmol) in CH_2Cl_2 (1.5 mL) with acetic anhydride (23.5 μ L, 0.25 mmol) and stir for 35 min. Add the reaction mixture to 0.2 N HCl (25 mL) and extract with CH_2Cl_2 (3 x 10 mL). Wash the combined organic layers with sat. NaHCO_3 (10 mL) and brine (10 mL), dry with MgSO_4 , and concentrate. Chromatograph on silica gel, eluting with 20:1 CH_2Cl_2 : CH_3OH , to obtain 127 mg of the desired product as a white foam (0.2 mmol, 82 %).
- 10 HRMS (FAB, $\text{M}+\text{H}^+$): m/e calc'd for $[\text{C}_{33}\text{H}_{38}\text{N}_3\text{O}_5\text{Cl}_2]^+$ 626.2189; found 626.2181.

Example 9

- 15 Treat a solution of the product from Example 7 (63.5 mg, 0.109 mmol) and $\text{CH}_3\text{CH}_2\text{I}$ (10 μ L, 0.13 mmol) in DMF (1.1 mL) with 50% KF-celite (40.7 mg) and stir at 60°C overnight. Add the reaction mixture to H_2O (50 mL) and extract with EtOAc (3 x 25 mL). Wash the combined organic layers with brine (25 mL), dry with MgSO_4 , and concentrate.
- 20 Chromatograph on silica gel, eluting with 10:1 CH_2Cl_2 : CH_3OH : NH_3 (aq), to obtain 42.7 mg of the product as a white foam (0.07 mmol, 64 %).
- HRMS (FAB, $\text{M}+\text{H}^+$): m/e calc'd for $[\text{C}_{33}\text{H}_{40}\text{N}_3\text{O}_4\text{Cl}_2]^+$ 612.2396; found 612.2405.

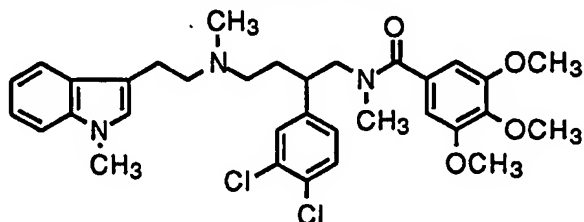
Example 9b

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Using the procedure of Example 9, substitute allyl bromide for $\text{CH}_3\text{CH}_2\text{I}$ to obtain the title compound.

HRMS (FAB, $\text{M}+\text{H}^+$): m/e calc'd for $[\text{C}_{34}\text{H}_{40}\text{N}_3\text{O}_4\text{Cl}_2]^+$ 624.2396; found 624.2385.

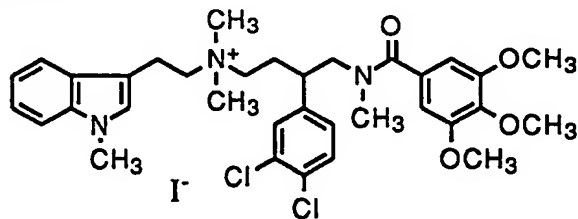
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Example 9c

Use a procedure similar to Example 9, substituting CH_3I (2 equivalents) for the alkylating agent and NaH for the base to obtain the title compound. HRMS (FAB, $\text{M}+\text{H}^+$): m/e calc'd for $[\text{C}_{33}\text{H}_{40}\text{N}_3\text{O}_4\text{Cl}_2]^+$ 612.2396; found 612.2407.

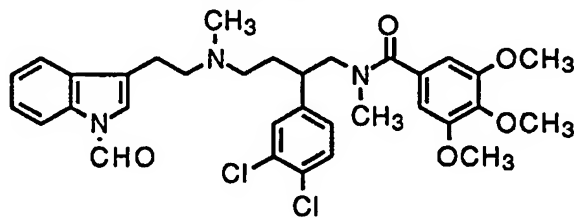
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The following compound is isolated as a side product from the previous reaction:



HRMS (FAB, M): m/e calc'd for $[\text{C}_{34}\text{H}_{42}\text{N}_3\text{O}_4\text{Cl}_2]^+$ 626.2552; found 626.2564.

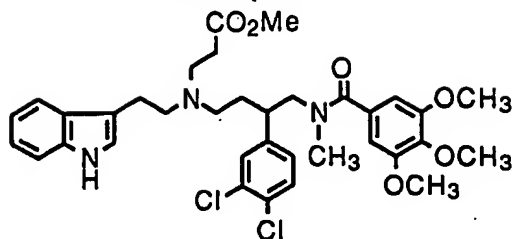
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Example 10

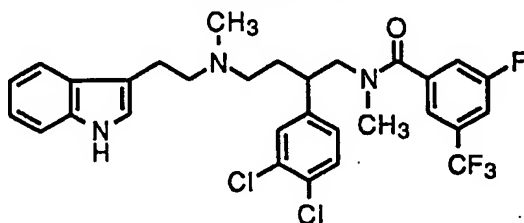
Treat the product of Example 2 (49.3 mg, 82 μmol) with formic acid (750 μL , 20 mmol) and acetic anhydride (30 mL, 0.32 mmol) and heat at 70°C for 3 days. Add the reaction mixture to H_2O (50 mL) and extract with CH_2Cl_2 (2 x 35 mL). Wash the combined organic layers with brine (25 mL), dry with MgSO_4 and concentrate. Chromatograph on silica gel, eluting with 20:1 to 10:1 $\text{CH}_2\text{Cl}_2:\text{CH}_3\text{OH}:\text{NH}_3$ (aq), to obtain 9.3 mg of the desired product as a white foam. HRMS (FAB, $\text{M}+\text{H}^+$): m/e calc'd for $[\text{C}_{33}\text{H}_{38}\text{N}_3\text{O}_5\text{Cl}_2]^+$ 626.2187; found 626.2189.

25

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Example 11

- Step 1:** Treat a solution of the product of Example 7 (115.1 mg, 0.197 mmol) in DMF (2.0 mL) with methyl acrylate (30 μ L, 0.33 mmol) and stir at 60°C overnight. Add additional methyl acrylate (20 μ L, 0.22 mmol) to the reaction mixture and stir at 60°C for 24 h. Add the reaction mixture to H₂O (50 mL) and extract with EtOAc (3 x 25 mL). Wash the combined organic layers with brine (25 mL), dry with MgSO₄, and concentrate. Chromatograph on silica gel, eluting with 25:1 CH₂Cl₂:CH₃OH, to obtain 66.9 mg of the desired product (0.10 mmol, 51 %). HRMS (FAB, M+H⁺): m/e calc'd for [C₃₅H₄₂N₃O₆Cl₂]⁺ 670.2451; found 670.2447.

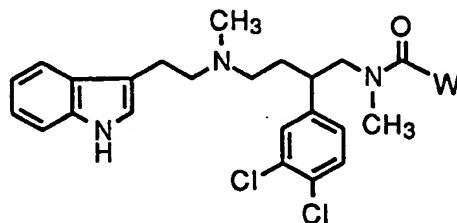
Example 12

- Step 1:** Heat a solution of the product of Example 1, step 4 (5.8 g, 17 mmol) in ethyl formate (100 mL) at reflux for 65 h and concentrate. Chromatograph on silica gel, eluting with 1:1 hexane:EtOAc, to obtain 4.4 g of the desired formamide as a colorless oil (12 mmol, 72 %).
- Step 2:** Treat a solution of the product of step 1 (4.2 g, 11.6 mmol) in Et₂O (25 mL) with BH₃.DMS (7.5 mL, 75 mmol) and stir at ambient temperature for 24 h. After concentrating the reaction mixture, quench the excess BH₃.DMS by the careful addition of CH₃OH. Add HCl-saturated CH₃OH to the resulting mixture, heat at 60°C for 1 h and concentrate to give 3.08 g of N-methyl-4-amino-3-(3,4-dichlorophenyl)-butanol.
- Step 3:** Treat the crude product of step 2 (3.08 g) with K₂CO₃ (3.02 g) and (BOC)₂O (3.43 g, 15.7 mmol) in CH₃OH (30 mL) for 2.5 h, filter and concentrate. Chromatograph on silica gel, eluting with 1:1 hexane:EtOAc, to obtain 2.50 g of the BOC-protected product (7.2 mmol, 62 % for 2 steps).

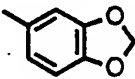
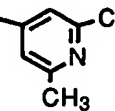
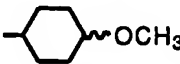
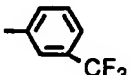
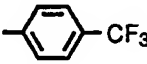
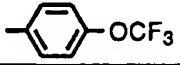
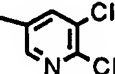
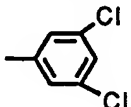
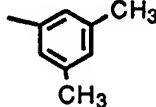
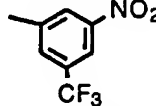
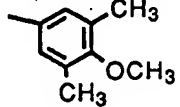
- 35 -

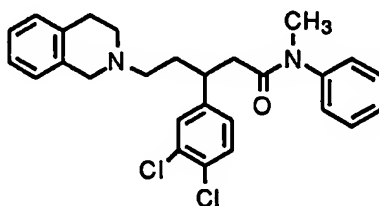
- Step 4:** To a -55°C solution of oxalyl chloride (0.94 mL, 10.8 mmol) in CH₂Cl₂ (20 mL), slowly add DMSO (1.0 mL, 14.1 mmol) followed by a solution of the alcohol from step 3 (2.50 g, 7.2 mmol) in CH₂Cl₂ (35 mL). After stirring for 30 min. at -55°C, add Et₃N (4.0 mL, 29 mmol) and stir for 2 h at -55°C. Add 20% sat. KHSO₄ (28 mL) and Et₂O (35 mL), warm to room temperature and stir for 30 min. Add to Et₂O (100 mL), remove the aqueous layer, wash with sat. NaHCO₃ (3 x 35 mL), H₂O (35 mL) and brine (35 mL). Dry the solution with MgSO₄ and concentrate to give 2.5 g of desired aldehyde product (quantitative yield).
- Step 5:** Treat a solution of the product of step 4 (2.50 g, 7.2 mmol), N-methyl tryptamine (13.8 mmol) and crushed 3A mol. sieves (2.0 g) in CF₃CH₂OH (25 mL) with NaCNBH₃ (1.75 g, 27.8 mmol). Stir for 1 h, add H₂O (250 mL) and extract with 2:1 CH₂Cl₂:Et₂O (3 x 150 mL). Wash the combined organic layers with brine (150 mL), dry with MgSO₄ and concentrate. Chromatograph the crude product on silica gel, eluting with 10:1:0.1 CH₂Cl₂:CH₃OH:NH₃ (aq.) to give 2.5 g of product as a white foam (5.0 mmol, 69 %).
- Step 6:** Treat the product of step 5 (4.5 g, 8.9 mmol) with trifluoroacetic acid (30 mL) for 2 h and concentrate. Suspend the residue in saturated NaHCO₃ and extract with CH₂Cl₂. Wash the combined organic layers with brine, dry with MgSO₄ and concentrate. Chromatograph on silica gel, eluting with 10 to 20 % NH₃ saturated CH₃OH:CH₂Cl₂, to obtain 2.0 g of the amino product as a white solid (4.9 mmol, 55 %).
- Step 7:** Convert the product from step 6 to the title compound using a procedure similar to that described in Example 2, Step 1, substituting 3,4,5-trimethoxybenzoic acid for 3-fluoro-5-trifluoromethylbenzoic acid. HRMS (FAB, M+H⁺): m/e calc'd for [C₃₀H₃₀N₃OC₂F₄]⁺ 594.1702; found 594.1702.

- Using the procedure of Example 12, substitute the appropriate acid in step 7 to obtain the compounds of the following structure, wherein W is as shown in the table:



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Ex.	W	HRMS (FAB, M+H ⁺): m/e calc'd	HRMS (FAB, M+H ⁺): m/e found
12a		552.1821	552.1816
12c		557.1642	557.1644
12d		544.2498	544.2499
12e		576.1796	576.1796
12f		576.1796	576.1792
12g		592.1745	592.1746
12h		577.1095	577.1087
12i			MS (FAB, M+H ⁺): m/e 576
12j		536.2235	536.2249
12k		621.1647	621.1646
12l		566.2341	566.2345

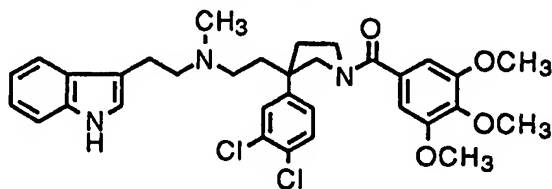
Example 13

Treat 3,4-dichloro-beta-(2-oxoethyl)-N-methyl-N-phenylbenzenepropanamide (0.53 g) in CH₃OH (35 mL) sequentially with
 5 molecular sieves 3A (5.5 g), isoquinoline HCl (0.33 g) and NaBH₃CN (0.4

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g). Stir the resulting mixture at room temperature for 20 hours, filter through a pad of celite and concentrate under reduced pressure. Partition the residue between 10% NH₄OH solution and CH₂Cl₂ (25 mL), separate the organic layer and extract the aqueous layer with CH₂Cl₂ (2X25 mL).

- 5 Dry the combined organic layers over MgSO₄, filter and concentrate under reduced pressure to give a crude oil (0.7 g). Chromatograph on silica gel, eluting with 2% CH₃OH/CH₂Cl₂ to obtain the title compound (0.27 g). Mass spectrum (FAB): 467.

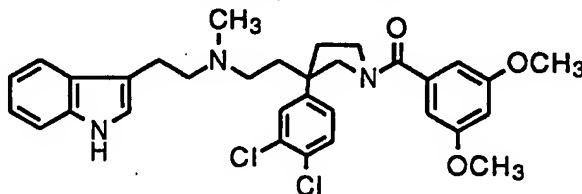
Example 14

10

Use 3-(3,4-dichlorophenyl)-3-(2-hydroxyethyl)-pyrrolidine in the procedure of Example 12, steps 3-7, substituting 3,4,5-trimethoxybenzoic acid for 3-fluoro-5-trimethylbenzoic acid to obtain the title compound.

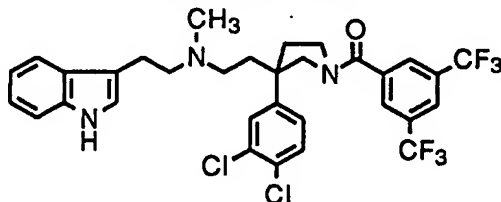
HRMS (FAB, M+H⁺): m/e calc'd for [C₃₃H₃₇N₃O₄Cl₂]⁺ 610.2239; found

15 610.2219.

Example 14a

Use the procedure of Example 14, substituting 3,5-dimethoxybenzoic acid for 3,4,5-trimethoxybenzoic acid to obtain the title compound.

20 HRMS (FAB, M+H⁺): m/e calc'd for [C₃₂H₃₆N₃O₃Cl₂]⁺ 580.2134; found 580.2116.

Example 14b

25 Using the procedure of Example 14, substitute 3,5-bis(trifluoromethyl)benzoic acid for 3,4,5-trimethoxybenzoic acid to obtain the title compound. HRMS (FAB, M+H⁺): m/e calc'd for [C₃₂H₃₀N₃OCl₂F₆]⁺ 656.1670; found 656.1663.

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The following formulations exemplify some of the dosage forms of this invention. In each, the term "active compound" refers to a compound of formula I.

EXAMPLE A

5

Tablets

<u>No.</u>	<u>Ingredient</u>	<u>mg/tablet</u>	<u>mg/tablet</u>
1	Active Compound	100	500
2	Lactose USP	122	113
3	Corn Starch, Food Grade, as a 10% paste in Purified Water	30	40
4	Corn Starch, Food Grade	45	40
5	Magnesium Stearate	<u>3</u>	<u>7</u>
	Total	300	700

Method of Manufacture

- Mix Item Nos. 1 and 2 in suitable mixer for 10-15 minutes. Granulate the mixture with Item No. 3. Mill the damp granules through a coarse screen (e.g., 1/4", 0.63 cm) if necessary. Dry the damp granules.
- 10 Screen the dried granules if necessary and mix with Item No. 4 and mix for 10-15 minutes. Add Item No. 5 and mix for 1-3 minutes. Compress the mixture to appropriate size and weight on a suitable tablet machine.

EXAMPLE BCapsules

<u>No.</u>	<u>Ingredient</u>	<u>mg/tablet</u>	<u>mg/tablet</u>
1	Active Compound	100	500
2	Lactose USP	106	123
3	Corn Starch, Food Grade	40	70
4	Magnesium Stearate NF	<u>4</u>	<u>7</u>
	Total	250	700

15 Method of Manufacture

Mix Item Nos. 1, 2 and 3 in a suitable blender for 10-15 minutes. Add Item No. 4 and mix for 1-3 minutes. Fill the mixture into suitable two-piece hard gelatin capsules on a suitable encapsulating machine.

EXAMPLE C

20

Sterile Powder for Injection

<u>Ingredient</u>	<u>mg/vial</u>	<u>mg/vial</u>
Active sterile powder	100	500

For reconstitution add sterile water for injection or bacteriostatic water for injection.

The *in vitro* and *in vivo* activity of the compounds of formula I can be determined by the following procedures.

In vitro procedure to identify NK₁ activity

- Test compounds are evaluated for their ability to inhibit the activity of the NK₁ agonist Substance P on the isolated guinea pig vas deferens. Freshly cut vas deferens are removed from male Hartley guinea pigs (230-350g) and suspended in 25 ml tissue baths containing Kreb's Henseleit solution warmed to 37°C and constantly aerated with 95% O₂ and 5% CO₂. Tissues are adjusted to 0.5 g and allowed to equilibrate for a period of 30 minutes. The vas deferens are exposed to an electrical field stimulation (Grass S48 Stimulator) every 60 seconds at an intensity that will cause the tissue to contract 80% of its maximum capacity. All responses are recorded isometrically by means of a Grass force displacement transducer (FT03) and Harvard electronic recorder.
- Substance P inhibits the electrical field stimulated-induced contractions of the guinea pig vas deferens. In unpaired studies, all tissues (control or drug treated) are exposed to cumulative concentrations of Substance P (1×10^{-10} M - 7×10^{-7} M). Single log-concentrations of the test compounds are given to separate tissues and allowed to equilibrate for 30 minutes before a Substance P concentration-response curve is generated. At least 5 separate tissues are used for each control and individual drug-concentration for every drug assay.

- Inhibition of the Substance P is demonstrated by a rightward shift of its concentration-response curve. These shifts are used to determine the pA₂ value, which is defined as the negative log of the molar concentration of the inhibitor which would require that twice as much agonist be used to elicit a chosen response. This value is used to determine relative antagonist potency.

Isolated Hamster Trachea NK₂ Assay

- General methodology and characterization of hamster trachea responses to neurokinin agonists as providing an NK₂ monoreceptor assay is found in C.A. Maggi, et al., *Eur. J. Pharmacol.* 166 (1989) 435 and J.L. Ellis, et al., *J. Pharm. Exp. Ther.* 267 (1993) 95.

- Continuous isometric tension monitoring is achieved with Grass FT-03 force displacement transducers connected to Buxco Electronics preamplifiers built into a Graphtec Linearrecorder Model WR 3310.

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Male Charles River LAK:LVG (SYR) hamsters, 100-200 g fed weight, are stunned by a sharp blow to the head, loss of corneal reflex is assured, the hamsters are sacrificed by thoractomy and cutting the heart. Cervical trachea segments are removed to room temperature Krebs buffer, pH 7.4, aerated with 95% O₂ - 5% CO₂ gas and cleaned of adhering tissue. The segments are cut into two 3-4 mm long ring segments. Tracheal rings are suspended from transducers and anchored in 15.0 ml water jacketed organ baths by means of stainless steel hooks and 6-0 silk. Baths are filled with Krebs buffer, pH 7.4, maintained at 37°C and continuously aerated with 95% O₂ - 5% CO₂ gas. Tracheal rings are placed under 1.0 g initial tension and allowed a 90 min equilibration period with four 1 µM NKA challenge, wash and recovery cycles at 20 min intervals. 30 min vehicle pretreatment is followed by cumulative additions of rising doses of NKA (3 nM - 1 µM final concentration, 5 min intervals between additions). The final NKA response is followed by a 15 min wash and recovery period. 30 min pretreatment with a test compound or its vehicle is followed by cumulative additions of rising doses of NKA (3 nM - 10 µM final concentration if necessary, 5 min intervals between additions). The final NKA response is followed by a 1 mM carbachol challenge to obtain a maximal tension response in each tissue.

Tissue responses to NKA are recorded as positive pen displacements over baseline and converted to grams tension by comparison to standard weights. Responses are normalized as a % of the maximal tissue tension. ED₅₀'s are calculated for NKA from the control and treated NKA dose responses and compared. Test compounds resulting in an agonist dose ratio ≥ 2 at a screening concentration of 1 µM (i.e. $pA_2 \geq 6.0$) are considered actives. Further dose response data is obtained for actives so that an apparent pA_2 estimate can be calculated. pA_2 is calculated either by estimation of K_i as described by Furchgott (where $pA_2 = -\log K_i$, R.F. Furchgott, *Pharm. Rev.* 7 [1995] 183) or by Schild Plot Analysis (O. Arunlakshana & H.O. Schild, *Br. J. Pharmacol.* 14[1959] 48) if the data is sufficient.

Effect of NK₁ Antagonists on Substance P-Induced Airway Microvascular Leakage in Guinea Pigs

Studies are performed on male Hartley guinea pigs ranging in weight from 400-650 g. The animals are given food and water *ad libitum*. The animals are anesthetized by intraperitoneal injection of dialurethane (containing 0.1 g/ml diallylbarbituric acid, 0.4 g/ml ethylurea

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and 0.4 g/ml urethane). The trachea is cannulated just below the larynx and the animals are ventilated ($V_T = 4$ ml, $f = 45$ breaths/min) with a Harvard rodent respirator. The jugular vein is cannulated for the injection of drugs.

- 5 The Evans blue dye technique (Danko, G. et al., Pharmacol. Commun., 1, 203-209, 1992) is used to measure airway microvascular leakage (AML). Evans blue (30 mg/kg) is injected intravenously, followed 1 min later by i.v. injection of substance P (10 μ g/kg). Five min later, the thorax is opened and a blunt-ended 13-gauge needle passed into the
10 aorta. An incision is made in the right atrium and blood is expelled by flushing 100 ml of saline through the aortic catheter. The lungs and trachea are removed en-bloc and the trachea and bronchi are then blotted dry with filter paper and weighed. Evans blue is extracted by incubation of the tissue at 37°C for 18 hr in 2 ml of formamide in stoppered tubes. The
15 absorbance of the formamide extracts of dye is measured at 620 nm. The amount of dye is calculated by interpolation from a standard curve of Evans blue in the range 0.5-10 μ g/ml in formamide. The dye concentration is expressed as ng dye per mg tissue wet weight. Test compounds were suspended in cyclodextran vehicle and given i.v. 5 min
20 before substance P.

Measurement of NK₂ Activity In Vivo

- Male Hartley guinea pigs (400-500 gm) with ad lib. access to food and water are anesthetized with an intraperitoneal injection of 0.9 ml/kg dialurethane (containing 0.1 g/m diallylbarbituric acid, 0.4 g/ml
25 ethylurea and 0.4 g/ml urethane). After induction of a surgical plane of anesthesia, tracheal, esophageal and jugular venous cannulae are implanted to facilitate mechanical respiration, measurement of esophageal pressure and administration of drugs, respectively.

- The guinea pigs are placed inside a whole body
30 plethysmograph and the catheters connected to outlet ports in the plethysmograph wall. Airflow is measured using a differential pressure transducer (Validyne, Northridge CA, model MP45-1, range ± 2 cmH₂O) which measures the pressure across a wire mesh screen that covers a 1 inch hole in the wall of the plethysmograph. The airflow signal is
35 electrically integrated to a signal proportional to volume. Transpulmonary pressure is measured as the pressure difference between the trachea and the esophagus using a differential pressure transducer (Validyne, Northridge, CA, model MP45-1, range ± 20 cm H₂O). The volume, airflow

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and transpulmonary pressure signals are monitored by means of a pulmonary analysis computer (Buxco Electronics, Sharon, CT, model 6) and used for the derivation of pulmonary resistance (R_L) and dynamic lung compliance ($C_{D_{\text{dyn}}}$).

5 Bronchoconstriction Due to NKA

Increasing iv doses of NKA are administered at half log (0.01-3 $\mu\text{g/kg}$) intervals allowing recovery to baseline pulmonary mechanics between each dose. Peak bronchoconstriction occurs within 30 seconds after each dose of agonist. The dose response is stopped
10 when $C_{D_{\text{dyn}}}$ is reduced 80-90% from baseline. One dose-response to NKA is performed in each animal. Test compounds are suspended in cyclodextran vehicle and given i.v. 5 min before the initiation of the NKA dose response.

For each animal, dose response curves to NKA are
15 constructed by plotting the percent increase in R_L or decrease in $C_{D_{\text{dyn}}}$ against log dose of agonist. The doses of NKA that increased R_L by 100% (R_L100) or decreased $C_{D_{\text{dyn}}}$ by 40% ($C_{D_{\text{dyn}}40}$) from baseline values are obtained by log-linear interpolation of the dose response curves.

Neurokinin Receptor Binding Assay(s)

20 Chinese Hamster ovary (CHO) cells transfected with the coding regions for the human neurokinin 1 (NK1) of the human neurokinin 2 (NK2) receptors are grown in Dulbecco's minimal essential medium supplemented with 10% fetal calf serum, 0.1 mM non-essential amino acids, 2 mM glutamine, 100units/ml of penicillin and streptomycin, and 0.8
25 mg of G418/ml at 37°C in a humidified atmosphere containing 5% CO₂.

Cells are detached from T-175 flasks with a sterile solution containing 5mM EDTA in phosphate buffered saline. Cells are harvested by centrifugation and washed in RPMI media at 40°C for 5 minutes. The pellet is resuspended in Tris-HCl (pH7.4) containing 1 μM phsphoramidon
30 and 4 $\mu\text{g/ml}$ of chymostatin at a cell density of 30×10^6 cells/ml. The suspension is then homogenized in a Brinkman Polytron (setting 5) for 30-45 seconds. The homogenate is centrifuged at 800 x g for 5 min at 4°C to collect unbroken cells and nuclei. The supernatant is centrifuged in a Sorvall RC5C at 19,000 rpm (44,00 x g) for 30 min at 4°C. The pellet is
35 resuspended, an aliquot is removed for a protein determination (BCA) and washed again. The resulting pellet is stored at -80°C.

To assay receptor binding, 50 μl of [³H]-Substance P (9-Sar, 11-Met [02]) (specific activity 41 Ci/mmol) (Dupont-NEN) (0.8 nM for the

- 43 -

NK-1 assay) or [³H]-Neurokinin A (specific activity 114 Ci/ mmole) (Zenca) (1.0 nM for the NK-2 assay) is added to tubes containing buffer (50 mM Tris-HCl (pH 7.4) with 1 mM MnCl₂ and 0.2% Bovine Serum Albumin) and either DMSO or test compound. Binding is initiated by the addition of 100 μl of membrane (10-20 μg) containing the human NK-1 or NK-2 receptor in a final volume of 200 μl. After 40 minutes at room temperature, the reaction is stopped by rapid filtration onto Whatman GF/C filters which have been presoaked in 0.3% polyethylenimine. Filters are washed 2 times with 3 ml of 50 mM Tris-HCl (pH 7.4). Filters are added to 6 mls of Ready-Safe liquid scintillation cocktail and quantified by liquid scintillation spectrometry in a LKB 1219 RackBeta counter. Non-specific binding is determined by the addition of either 1 μM of CP-99994 (NK-1) or 1 μM SR-48968 (NK-2) (both synthesized by the chemistry department of Schering-Plough Research Institute). IC₅₀ values are determined from competition binding curves and K_i values are determined according to Cheng and Prusoff using the experimentally determined value of 0.8 nM for the NK-1 receptor and 2.4 nM for the NK-2 receptor.

NK₃ activity is determined by following a procedure similar to that described in the literature, e.g., *Molecular Pharmacol.*, 48 (1995), p. 711-716.

% Inhibition is the difference between the percent of maximum specific binding (MSB) and 100%. The percent of MSB is defined by the following equation, wherein "dpm" is disintegrations per minute:

$$\% \text{ MSB} = \frac{(\text{dpm of unknown}) - (\text{dpm of nonspecific binding})}{(\text{dpm of total binding}) - (\text{dpm of nonspecific binding})} \times 100$$

It will be recognized that compounds of formula I exhibit NK₁, NK₂ and/or NK₃ antagonist activity to varying degrees, e.g., certain compounds have strong NK₁ antagonist activity, but weaker NK₂ and NK₃ antagonist activity, while others are strong NK₂ antagonists, but weaker NK₁ and NK₃ antagonists. While compounds with approximate equipotency are preferred, it is also within the scope of this invention to use compounds of with unequal NK₁/NK₂/NK₃ antagonist activity when clinically appropriate.

Using test procedures described above, the following data (% inhibition or K_i) were obtained for preferred and/or representative compounds of formula I:

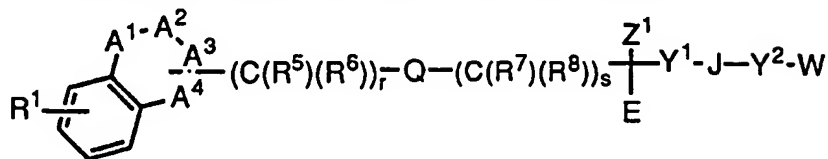
Ex.	% Inhibition NK ₁ (1μM dose)	Ki (NK ₁) (nM)	% Inhibition NK ₂ (1μM dose)	Ki (NK ₂) (nM)	Ki (NK ₃) (nM)
7c	--	12	100	10	--
7d	--	233	--	30	--
9b	--	23	--	250	--
12a	69	165	96	59	--
12k	--	30	--	39	--
14a	--	233	--	30	--
2	81	25	87	33	448

Compounds of the present invention exhibit a range of activity: percent inhibition at a dosage of 1μM ranges from about 0 to about 100% inhibition of NK₁ and/or about 0 to about 100% inhibition of NK₂. Preferred are compounds having a Ki ≤100nM for the NK₁ receptor.

- 5 Also preferred are compounds having a Ki ≤100nM for the NK₂ receptor. Another group of preferred compounds are those having a Ki ≤100nM for each of the NK₁ and NK₂ receptors.

We claim:

1. A compound represented by the structural formula



- 5 or a pharmaceutically acceptable salt thereof, wherein:

A¹, A², A³ and A⁴ are independently selected from the group consisting of -(C(R²)(R¹⁰))- , -(C(R^{2c})(R¹⁰))- , -(C(R²))= , -(C(R^{2c}))= , -NR¹⁶- , -N= , -O- , -S(O)_e- , -C(O)- and a bond, wherein A¹, A², A³ and A⁴, together with the carbon atoms to which they are attached, form a 5- or 6-
 10 membered ring, and wherein two adjacent A groups are selected from the group consisting of the following combinations:

- (C(R²)(R¹⁰))- (C(R²)(R¹⁰))- ;

- (C(R²)(R¹⁰))- (C(R^{2c}))= ;

- (C(R^{2c})(R¹⁰))- NR¹⁶- ;

15 - (C(R^{2c})(R¹⁰))- N= ;

- (C(R^{2c})(R¹⁰))- O- ;

- (C(R^{2c})(R¹⁰))- S(O)_e- ;

- (C(R²)(R¹⁰))- C(O)- ;

- (C(R²))= (C(R²))- ;

20 - (C(R²))= N- ;

= (C(R²))- (C(R²))= ;

= (C(R²))- NR¹⁶- ;

= (C(R²))- N= ;

= (C(R²))- O- ;

25 = (C(R²))- S(O)_e- ;

= (C(R²))- C(O)- ;

- NR¹⁶- N= ;

- NR¹⁶- S(O)_e- ;

- NR¹⁶- C(O)- ;

30 - N= N- ;

= N- N= ;

= N- O- ;

= N- S(O)_e- ;

= N- C(O)- ;

35 - O- S(O)_e- ; and

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-O-C(O)-;

provided that three adjacent A groups do not represent -C(O)-O-C(O)-, -S(O)-O-C(O)- or -S(O)-O-S(O)-, and provided that when an aromatic nitrogen is present in the ring formed by A¹, A², A³ and A⁴, the N-oxide can be formed;

E is R³-aryl or R³-heteroaryl;

W is R⁴-cycloalkyl, R⁴-aryl, R⁴-heterocycloalkyl or R⁴-heteroaryl;

R¹, R³ and R⁴ are independently 1-3 substituents

independently selected from the group consisting of H, halogeno, C₁-C₆ alkyl, -CF₃, -C₂F₅, -OR¹¹, -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), -N(R¹¹)(R¹²), -N(R¹¹)COR¹², -SH, -S(O)_eR¹³, -OC(O)R¹¹, -OC(O)N(R¹¹)(R¹²), -NR¹¹CO₂R¹³, -NR¹¹C(O)N(R¹²)(R¹⁴), R¹⁵-phenyl, R¹⁵-benzyl, -NO₂, -NR¹¹S(O)₂R¹³ and -S(O)₂NR¹¹R¹²; or adjacent R¹, R³ or R⁴ substituents can form a -O-CH₂-O- group;

R² is independently selected from the group consisting of R^{2a}, R^{2b} or R¹⁰; and R^{2c} is independently selected from the group consisting of R^{2a} and R¹⁰; wherein R^{2a} is selected from the group consisting of -CF₃, -C₂F₅, -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), R¹⁵-phenyl and R¹⁵-benzyl; and R^{2b} is selected from the group consisting of halogeno, -OR¹¹, -NO₂, -N(R¹¹)(R¹²), -N(R¹¹)COR¹², -SH, -S(O)_eR¹³, -OC(O)R¹¹, -OC(O)N(R¹¹)(R¹²), -NR¹¹CO₂R¹³, -NR¹¹C(O)N(R¹²)(R¹⁴), -NR¹¹S(O)₂R¹³ and -S(O)₂NR¹¹R¹²; provided that from any combination of ring members A¹, A², A³ and A⁴ comprising R² and R^{2c}, no more than one substituent can be selected from R^{2a} and no more than one substituent can be selected from R^{2b};

R⁵, R⁷, R⁹, R¹¹, R¹² and R¹⁴ are independently selected from the group consisting of H, C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃ and -C₂F₅;

R⁶ and R⁸ are independently selected from the group consisting of R⁵, -(C(R⁹)(R¹⁰))_n-OR¹¹, -(C(R⁹)(R¹⁰))_n-NR¹¹R¹², -(C(R⁹)(R¹⁰))_n-SH, -(C(R⁹)(R¹⁰))_n-S(O)_eR¹³, -(C(R⁹)(R¹⁰))_n-CO₂R¹¹, -(C(R⁹)(R¹⁰))_n-OC(O)R¹¹, -(C(R⁹)(R¹⁰))_n-CONR¹¹R¹², -(C(R⁹)(R¹⁰))_n-COR¹¹ and -(C(R⁹)(R¹⁰))_n-NR¹¹C(O)R¹², provided that when Q is a heteroatom, R⁶ and R⁸ cannot be -OR¹¹, -OC(O)R¹¹, -N(R¹¹)COR¹², -NR¹¹R¹², -SH or -S(O)_eR¹³ on adjacent carbon atoms;

R¹⁰ is independently selected from the group consisting of hydrogen and C₁-C₆ alkyl;

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R¹³ is independently selected from the group consisting of C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃ and -C₂F₅;

R¹⁵ is hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy or halogeno;

Q is a bond, -C(O)-, -NR¹⁷-, -(C(R⁹)(R¹⁰))-, -O-, -S(O)_e-,

5 -C(X)NR¹¹-, -N(R¹¹)C(X)-, -N(R¹¹)SO₂-, -SO₂N(R¹¹)- or -N⁺(R¹¹)(R¹⁷)-;

R¹⁶ and R¹⁷ are independently selected from the group consisting of hydrogen, C₁-C₆ alkyl, -S(O)_eR¹³, -COR¹¹, -(CH₂)_m-CO₂R¹³, -CONR¹¹R¹², C₂-C₆ alkenyl, -R¹⁵-phenyl and R¹⁵-benzyl;

X is =O, =S or =N(R¹²);

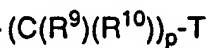
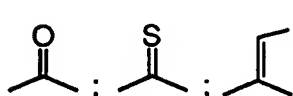
10 Y¹ is -(C(R⁹)(R¹⁰))_m-, -G-(C(R⁹)(R¹⁰))_m- or -(C(R⁹)(R¹⁰))_m-G-;

G is





, provided that when m is 0, R² is H, C₁-C₆ alkyl, -CF₃, -C₂F₅,

-COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), R¹⁵-phenyl or R¹⁵-benzyl;



, provided that when p is 0, T is

not OH or -NR¹¹R¹²;  , provided that when r is 1,

T is not OR¹¹ or -NR¹¹R¹²; or  , provided T is not

-OR¹¹, -N(R¹¹)(R¹²), -S(O)_eR¹³, -NR¹¹CO₂R¹³, -NR¹¹COR¹², -NR¹¹CON(R¹²)(R¹⁴) or -OC(O)N(R¹¹)(R¹²);

15 T is H, R¹⁵-aryl, R¹⁵-heterocycloalkyl, R¹⁵-heteroaryl, R¹⁵-cycloalkyl, -OR¹¹, -N(R¹¹)(R¹²), -COR¹¹, -CO₂R¹¹, -CON(R¹¹)(R¹²), -S(O)_eR¹³, -NR¹¹CO₂R¹³, -NR¹¹COR¹², -NR¹¹CON(R¹²)(R¹⁴) or -OC(O)N(R¹¹)(R¹²);

20 J is a bond, -S(O)_e-, -O- or -N(Z²)-, -N(Z²)C(O)- or -N(Z²)C(S)-; and when G is -C(R²)H-, J can also be -N(Z²)C(O)O- or -OC(O)N(Z²)-;

Y² is -(C(R⁹)(R¹⁰))_m-;

25 Z¹ is H, C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃, -C₂F₅, -NR¹¹R¹², -OR¹¹ or SR¹¹; Z² is H, C₁-C₆ alkyl, R¹⁵-phenyl, R¹⁵-benzyl, -CF₃ or -C₂F₅; provided that when Y¹ is -(C(R⁹)(R¹⁰))_m- and m is 0, Z¹ is not -NR¹¹R¹², -OR¹¹ or -SR¹¹; or Z¹ and Z² together are -(C(R⁹)(R¹⁰))_u-;

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wherein u is 1 to 4, and wherein with the atoms to which they are attached, form a 4 to 8 membered ring;

e and n are independently 0, 1 or 2;

m and p are independently 0, 1, 2 or 3; and

5 r and s are independently 1, 2, 3 or 4.

2. A compound of claim 1 wherein E is R³-phenyl.

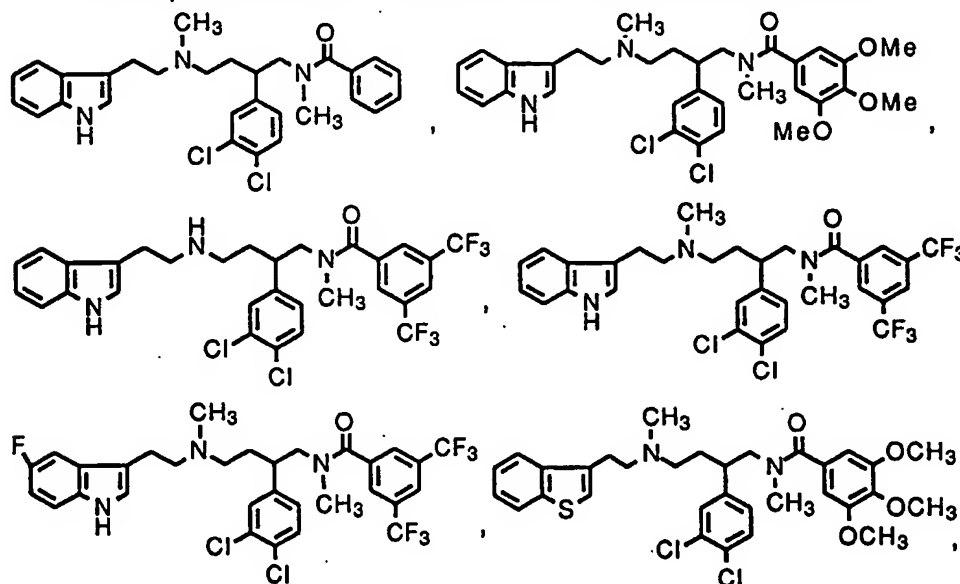
3. A compound of claim 1 or 2 wherein Z¹ is hydrogen, or Z¹ and Z² together are ethylene or propylene, and with the atoms to which they are attached form a 5- or 6-membered ring.

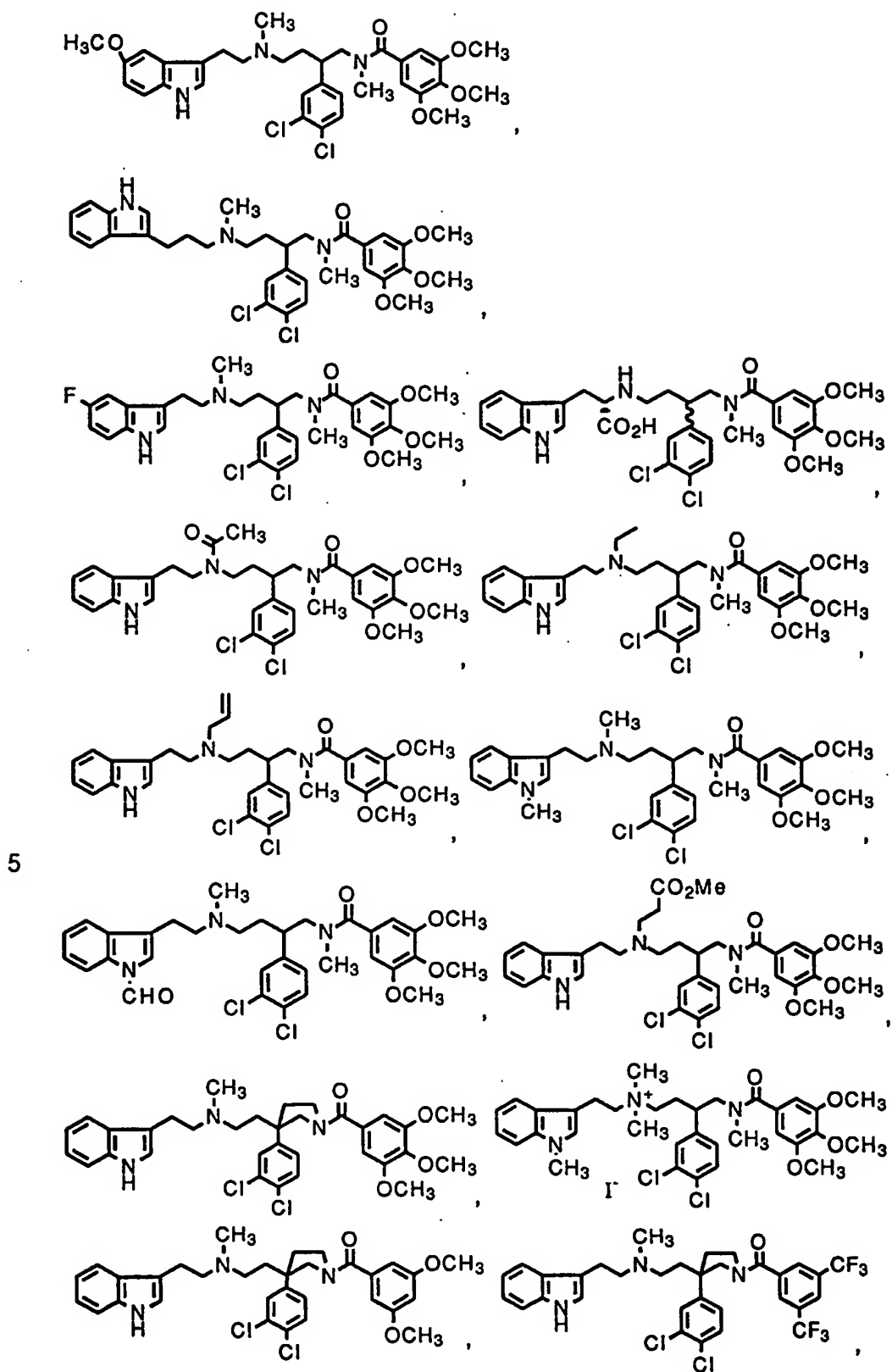
4. A compound of any of claims 1, 2 or 3 wherein Q is -NR¹⁷-, -O- or -S(O)_e-.

5. A compound of any of claims 1, 2, 3 or 4 wherein J is -O-, -N(Z²)- or -N(Z²)C(O)-.

6. A compound of any of claims 1, 2, 3, 4 or 5 wherein A¹, A², A³ and A⁴ comprise an indolyl ring.

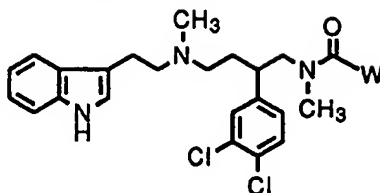
7. A compound of claim 1 selected from the group consisting of



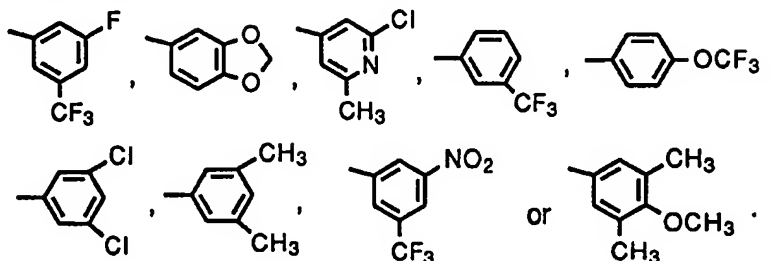


- 50 -

and a compound of the formula



wherein W is



5

8. A pharmaceutical composition comprising an effective amount of a compound of any of claims 1, 2, 3, 4, 5, 6 or 7 in a pharmaceutically acceptable carrier.
- 10 9. The use of a compound of any of claims 1, 2, 3, 4, 5, 6 or 7 for the preparation of a medicament for the treatment of asthma, cough, bronchospasm, central nervous system diseases, inflammatory diseases and gastrointestinal disorders.
- 15 10. A method of treating asthma, cough, bronchospasm, central nervous system diseases, inflammatory diseases and gastrointestinal disorders comprising administering an effective amount of a compound of any of claims 1, 2, 3, 4, 5, 6 or 7 to a mammal in need of such treatment.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 96/07960

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D209/16 A61K31/40 C07D333/58 C07D403/12 C07D209/20
C07D405/12 C07D401/12 C07D215/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US,A,5 350 852 (SANOFI) 27 September 1994 cited in the application see claims ---	1,8,9
A	EP,A,0 591 040 (ELF SANOFI) 6 April 1994 see claims ---	1,8,9
A	WO,A,93 18023 (MERCK SHARP & DOHME LTD.) 16 September 1993 see claims -----	1,8,9

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

8 August 1996

Date of mailing of the international search report

20.08.96

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
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Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 96/07960

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claim 10 is directed to a method of treatment of (diagnostic method practised on) the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/US 96/07960

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A-5350852	27-09-94	FR-A- 2666335	06-03-92
		FR-A- 2678267	31-12-92
		AU-B- 657272	09-03-95
		AU-B- 8354291	12-03-92
		CA-A- 2050639	06-03-92
		EP-A- 0474561	11-03-92
		IL-A- 99320	31-07-95
		JP-A- 4261155	17-09-92
		LT-A,B 585	27-12-94
		LV-B- 10606	20-04-96
		NO-B- 177226	02-05-95
		NZ-A- 239661	27-06-94
		PL-B- 167994	30-12-95
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EP-A-591040	06-04-94	FR-A- 2696178	01-04-94
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		CA-A- 2107432	31-03-94
		CN-A- 1088582	29-06-94
		CZ-A- 9302035	13-04-94
		FI-A- 934273	31-03-94
		HU-A- 65759	28-07-94
		JP-A- 6211850	02-08-94
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		SI-A- 9300513	30-06-94
		ZA-A- 9307276	25-04-94
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		CA-A- 2129145	16-09-93
		EP-A- 0629196	21-12-94
		JP-T- 7505147	08-06-95
		US-A- 5328927	12-07-94
		US-A- 5514718	07-05-96